

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:08 ; Search time 178 Seconds

(without alignment)
1668.573 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLESLSLGRGAPTV.....HKLNKTHDMLNENHEKLSLS 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1612378 seqs, 512079187 residues

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: uniprot_spot:*
2: uniprot_trembl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3055	100.0	580	1 MBD_HUMAN	095243 homo sapien
2	1821.5	59.6	554	1 MBD_MOUSE	0922d7 mus musculi
3	880.5	28.8	416	2 Q919F1	0919f1 gallus galli
4	344	11.3	419	2 Q9SFC1	09sfc1 arabidopsis
5	338	11.1	407	2 Q8AWT3	08awt3 arabidopsis
6	238.5	7.8	524	2 Q71T77	071t77 brachydanio
7	220.5	7.2	467	2 Q9YGC6	09ygc6 xenopus lae
8	213	7.0	682	2 Q7SCQ2	07scq2 neosporea
9	208.5	6.8	486	1 MEC2_HUMAN	P51608 homo sapien
10	208.5	6.8	498	2 Q6OH99	06oh99 homo sapien
11	208.5	6.8	516	2 Q7Z384	07z384 homo sapien
12	207.5	6.6	486	1 MEC2_MACFA	0951g8 macaca fasci
13	202.5	6.6	344	2 Q42403	042403 gallus galli
14	201	6.6	492	1 MEC2_RAT	000566 rattus norv
15	198.5	6.5	484	1 MEC2_MOUSE	0922d6 mus musculi
16	156	5.1	2473	2 Q7RN41	07rn41 plasmodium
17	149.5	4.9	2212	2 Q81T76	081t76 plasmodium
18	148.5	4.9	352	2 Q8L576	08l576 arabidopsis
19	148.5	4.9	1679	2 Q7RCP2	07rcp2 plasmodium
20	146	4.8	702	2 Q9SFC2	09sfc2 arabidopsis
21	146	4.8	1643	2 Q8HZQ1	08hzq1 pan troglod
22	145	4.7	267	2 Q8AYT1	08ayt1 xenopus lae
23	145	4.7	1301	1 SAC3_YEAST	P46674 saccharomyc
24	145	4.7	3418	1 BRG2_HUMAN	P51587 homo sapien
25	144	4.7	283	2 Q8AYP2	08ayp2 xenopus lae
26	143.5	4.7	782	2 Q25875	025875 plasmodium
27	141.5	4.6	782	2 Q25730	025730 plasmodium
28	141.5	4.6	782	2 Q26007	026007 plasmodium
29	140.5	4.6	782	2 Q9U414	09u414 plasmodium
30	140.5	4.6	782	2 Q9U430	09u430 plasmodium
31	140.5	4.6	881	2 Q7RK82	07rk82 plasmodium

32	140.5	4.6	1030	2 Q92AD9	092ad9 meriones sh
33	138.5	4.5	500	1 GAR2_SCHPO	P41891 schizosacch
34	138.5	4.5	782	2 Q9U429	09u429 plasmodium
35	138.5	4.5	1047	2 Q7RPV9	07rpv9 plasmodium
36	137.5	4.5	326	2 Q869D5	0869d5 branchiost
37	137.5	4.5	412	2 Q6MFI1	06mfi1 neosporea
38	137.5	4.5	782	2 Q26104	026104 plasmodium
39	137.5	4.5	2209	2 Q9U0G6	09u0g6 plasmodium
40	137	4.5	400	2 Q8LX06	08lx06 zea mays (m
41	137	4.5	560	1 SET4_YEAST	P42948 saccharomyc
42	137	4.5	707	2 Q7RKH5	07rkh5 plasmodium
43	137	4.5	1713	2 Q7TTS0	07tts0 mus musculi
44	137	4.5	1813	2 Q75X83	075x83 helicobacte
45	136.5	4.5	598	1 CYL1_HUMAN	P35663 homo sapien

ALIGNMENTS

RESULT 1
MED4_HUMAN STANDARD; PRT; 580 AA.
AC Q95243; Q724T3; Q96F09;
DT 25-OCT-2004 (Rel. 45, Created)
DT 25-OCT-2004 (Rel. 45, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-Cpg binding protein 4 (BC 3.2.2.-) (Methyl-Cpg binding domain
protein 4) (Methyl-Cpg binding endonuclease 1) (Mismatch-specific DNA
N-glycosylase).
GN Name=MBD4; Synonyms=MBD1;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP MEDLINE=98449942; PubMed=9774669;
RX Hendrich B., Bird A.;
RT "Identification and characterization of a family of mammalian methyl-
Cpg binding proteins.";
RL Mol. Cell. Biol. 18:6538-6547(1998).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RX MEDLINE=99373255; PubMed=10441743;
RA Hendrich B., Abbott C., McQueen H., Chambers D., Cross S.H., Bird A.;
RT "Genomic structure and chromosomal mapping of the murine and human
mbd1, mbd2, mbd3, and mbd4 genes.";
RL Mamm. Genome 10:906-912(1999).
RN [3]
RP SEQUENCE FROM N.A. (ISOFORM 1), FUNCTION AND INTERACTION WITH MLH1.
RX TISSUE=Petal brain;
RC MEDLINE=99199294; PubMed=10097147; DOI=10.1073/pnas.96.7.3969;
RA Bellacosa A., Cicchilitti L., Schepis F., Riccio A., Yeung A.T.,
RA Matsumoto Y., Golemis E.A., Genuardi M., Neri G.;
RT "MBD1, a novel human methyl-Cpg-binding endonuclease, interacts with
Proc. Natl. Acad. Sci. U.S.A. 96:3969-3974(1999).
RN [4]
RP SEQUENCE FROM N.A. (ISOFORM 3).
RX TISSUE=Lung;
RA Guo J.H., Chen L., Yu L.;
RT Submitted (Jul-2002) to the EMBL/Genbank/DBJ databases.
RN [5]
RP SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS SER-273; PRO-342; LYS-346
AND HIS-568.
RA Rieder M.J., Braun A.C., Montoya M.A., Chung M.-W., Nguyen C.P.,
RA Nguyen D.A., Livingston R.J., Poel C.L., Robertson P.D.,
RA Schackwitz W.S., Sherwood J.K., Wilczek L.A., Nickerson D.A.;
RT "NIHES-SNP, environmental genome project, NIHES ES15478, Department
of Genome Sciences, Seattle, WA (URL: <http://egp.gs.washington.edu>).";
RL Submitted (MAR-2002) to the EMBL/Genbank/DBJ databases.
RN [6]
RP SEQUENCE FROM N.A. (ISOFORM 2).

RA Ebert L., Schick M., Neubert P., Schatten R., Henze S., Korn B.;
RT "Cloning of human full open reading frames in Gateway(TM) system entry
RT vector (pDONR201)." ;
RL Submitted (MAY-2004) to the EMBL/GenBank/DBJ databases.
RN [7]
RP SEQUENCE FROM N.A. (ISOFORM 2) .
RC TISSUE=Lung;
RX MEDLINE=23388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Straubeberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shemmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Ditschenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldi M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carrinci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
RA Bosak S.A., McEwen P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman V.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butcherfield Y.S.N., Krzywinski M.I., Skalska U., Smalins D.E.,
RA Schermer A., Schein J.R., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences." ;
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
RN [8]
RP FUNCTION.
RX PubMed=10930409; DOI=10.1074/jbc.M004535200;
RA Petronzelli F., Riccio A., Markham G.D., Seeholzer S.H., Stoerker J.,
RA Genuardi M., Yeung A.T., Matsumoto Y., Bellacosa A.;
RT "Biaphasic kinetics of the human DNA repair protein MBD1 (MBD4), a
RT mismatch-specific DNA N-glycosylase." ;
RL J. Biol. Chem. 275:32422-32429 (2000).
RN [9]
RP INTERACTION WITH FADD.
RX PubMed=12702765; DOI=10.1073/pnas.0431215100;
RA Screation R.A., Kieselring S., Sansom O.J., Miller C.B., Maddison K.,
RA Bird A., Clarke A.R., Fritsch S.M.;
RT "Fas-associated death domain protein interacts with methyl-CpG binding
RT domain protein 4: a potential link between genome surveillance and
RT apoptosis." ;
RL Proc. Natl. Acad. Sci. U.S.A. 100:5211-5216 (2003).
CC -1- FUNCTION: Mismatch-specific DNA N-glycosylase involved in DNA
CC repair. Has thymine glycosylase activity and is specific for G:T
CC mismatches within methylated and unmethylated CpG sites. Can also
CC remove uracil or 5-fluorouracil in G:U mismatches. Has no lyase
CC activity. Was first identified as methyl-CpG-binding protein.
CC -1- SUBUNIT: Interacts with MHL1.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=3;
CC Name=1;
CC IsoId=O95243-1; Sequence=Displayed;
CC Name=2;
CC IsoId=O95243-2; Sequence=VSP_010816;
CC Note=No experimental confirmation available;
CC Name=3;
CC IsoId=O95243-3; Sequence=VSP_010817, VSP_010818;
CC Note=No experimental confirmation available;
CC -1- SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.
CC -----
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CC -----
CC EMBL; AF072250; AAC68879.1; -.

DR EMBL; AF120999; AAD50374.1; -.
DR EMBL; AF120997; AAD50374.1; JOINED.
DR EMBL; AF120998; AAD50374.1; JOINED.
DR EMBL; AF114784; AAD22195.1; -.
DR EMBL; AF532602; AAP97358.1; -.
DR EMBL; AF494057; AAM00008.1; -.
DR EMBL; CR450305; CAG29301.1; -.
DR EMBL; BC011752; AAH11752.1; -.
DR HSSP; Q922D7; INGN.
DR InrAct; O95243; -.
DR GeneW; HGNC:6919; MBD4.
DR H-InVDB; HIX0003669; -.
DR Reactome; O95243; -.
DR MIM; 603574; -.
DR GO; GO:0005634; C:nucleus; TAS.
DR GO; GO:0004520; F:endodeoxyribonuclease activity; TAS.
DR GO; GO:0003696; F:satellite DNA binding; TAS.
DR GO; GO:0006281; P:DNA repair; TAS.
DR InterPro; IPR003265; Endo 3c.
DR InterPro; IPR01739; Methyl-CpG_bind.
DR Pfam; PF00730; Hbh-GPD; 1.
DR Pfam; PF01429; MBD; 1.
DR SMART; SM00391; MBD; 1.
DR PROSITE; PS50982; MBD; 1.
DR KW Alternative splicing; DNA repair; DNA-binding; Hydrolase;
KW Nuclear protein; Polymorphism.
FT DOMAIN 76 148
FT MBD.
FT DOMAIN 461 524
FT Hbh-GPD.
FT ACT_SITE 560 560
FT Missing (in isoform 2).
FT VARSPLIC 395 400
FT /FTId=VSP_010816.
FT VARSPLIC 539 540
FT /FTId=VSP_010817.
FT VARSPLIC 541 580
FT /FTId=VSP_010818.
FT VARSPLIC 273 273
FT /FTId=VAR_019357.
FT VARIANT 273 273
FT A -> S (in dbSNP:103442) .
FT /FTId=VAR_019357.
FT VARIANT 342 342
FT S -> P (in dbSNP:2307289) .
FT /FTId=VAR_019514.
FT VARIANT 346 346
FT E -> K (in dbSNP:140693) .
FT /FTId=VAR_019358.
FT VARIANT 358 358
FT I -> T (in dbSNP:2307298) .
FT /FTId=VAR_019359.
FT VARIANT 568 568
FT D -> H (in dbSNP:2307293) .
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SQ SEQUENCE 580 AA; 66050 MW; BF16FB21A348B8E5F CRC64;
Query Match 100.0%; Score 3055; DB 1; Length 580;
Best Local Similarity 100.0%; Pred. No. 5,3e-178; Mismatches 0; Gaps 0;
Matches 580; Conservative 0; Indels 0;
QY 1 MGTGLSTSLGDRGAAPVTYSSRLVDPDPNDLRKEDVAMELERVGEDEEQMMIKRSSE 60
DB 1 MGTGLSTSLGDRGAAPVTYSSRLVDPDPNDLRKEDVAMELERVGEDEEQMMIKRSSE 60
QY 61 CNPILQEPPIASAOAGTAGTCRCRSVPCGWERVVKQQLFKGTAGRFVYFISPGGLKFRS 120
DB 61 CNPILQEPPIASAOAGTAGTCRCRSVPCGWERVVKQQLFKGTAGRFVYFISPGGLKFRS 120
QY 121 KSSLANTLHNKGESLKPEDPDFVLSKRGIKSRKYKCSMAALTSHLQNSNNNSNMMLRT 180
DB 121 KSSLANTLHNKGESLKPEDPDFVLSKRGIKSRKYKCSMAALTSHLQNSNNNSNMMLRT 180
QY 181 RSKCKDVFPMPSSSSSELQSRGLSNFTSTHLLKDEGVDDVFRKYRKPGRKVTLLKG 240
DB 181 RSKCKDVFPMPSSSSSELQSRGLSNFTSTHLLKDEGVDDVFRKYRKPGRKVTLLKG 240
QY 241 IPIKTKKGRKSCSGVQSDSKRESVCNKADASEEVAQXSQLDRVTCISDAGACETL 300
DB 241 IPIKTKKGRKSCSGVQSDSKRESVCNKADASEEVAQXSQLDRVTCISDAGACETL 300
QY 241 IPIKTKKGRKSCSGVQSDSKRESVCNKADASEEVAQXSQLDRVTCISDAGACETL 300
DB 241 IPIKTKKGRKSCSGVQSDSKRESVCNKADASEEVAQXSQLDRVTCISDAGACETL 300

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OY 301 SYTSENSLVKKKERSLSGSGNFCSEOKTSGTINKFCGSAKSEHNEKEDFLESEELGT 360
DB 301 SYTSENSLVKKKERSLSGSGNFCSEOKTSGTINKFCGSAKSEHNEKEDFLESEELGT 360
OY 361 KVEYVERKEHLHTDILKXGSEMDNNSCPFRKDPFTEKI FOEDPTI PRTOIERRKTSLYFSS 420
DB 361 KVEYVERKEHLHTDILKXGSEMDNNSCPFRKDPFTEKI FOEDPTI PRTOIERRKTSLYFSS 420
OY 421 KYNKALSPPRRKAFKWTTPRRSPNLVQETL FHPDWKLLIATIFLANTSGMAIPVLWK 480
DB 421 KYNKALSPPRRKAFKWTTPRRSPNLVQETL FHPDWKLLIATIFLANTSGMAIPVLWK 480
OY 481 FLEKTPSAEAVATAWRDVSSELKVLGLYDLAKTIKVSDEYTLKQWKYPIELHIGIKY 540
DB 481 FLEKTPSAEAVATAWRDVSSELKVLGLYDLAKTIKVSDEYTLKQWKYPIELHIGIKY 540
OY 541 GNDSYRIFCVNEMKOVHPEDHLKYNKHPDLNENHEKLSLS 580
DB 541 GNDSYRIFCVNEMKOVHPEDHLKYNKHPDLNENHEKLSLS 580

RESULT 2
MED4_MOUSE STRAND: PRT; 554 AA.
ID MED4_MOUSE Q922D7; Q792D2; Q8R3R3;
AC 25-OCT-2004 (Rel. 45, Created)
DT 25-OCT-2004 (Rel. 45, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-Cpg binding protein 4 (BC 3.2.2.-) (Methyl-Cpg binding domain
  protein Mbd4) (Mismatch-specific DNA N-glycosylase).
GN Name=Mbd4;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A., FUNCTION, AND SUBCELLULAR LOCATION.
RX MEDLINE=98449942; PubMed=9774669;
RT Hendrich B., Bird A.;
RT "Identification and characterization of a family of mammalian methyl-
  Cpg binding proteins.";
RT Mol. Cell. Biol. 18:6538-6547 (1998).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=129;
RC MEDLINE=9937325; PubMed=10441743;
RT Hendrich B., Abbott C., McQueen H., Chambers D., Cross S.H., Bird A.;
RT "Genomic structure and chromosomal mapping of the murine and human
  Mamm. Genome 10:906-912 (1999).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Klausner R.L., Feingold E.A., Grouse L.H., Derge J.G., Schuler G.D.,
RA Altschul S.F., Collins F.S., Wagner L., Shennan C.M., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Heist F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Donald M.F., Casavant T.L., Schetz T.E.,
RA Brownstein M.J., Ueda T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosak S.A., McMan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Huliy S.W.,
RA Villalon D.K., Muzny K.C., Sodergren B.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalins D.E.,
RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
  and mouse cDNA sequences.";
RT Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
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RN [4]
RP X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF 411-554.
RX PubMed12456671; DOI=10.1074/jbc.M210848200;
RA Wu P., Qiu C., Schall A., Zhang X., Bhagwat A.S., Cheng X.,
RT "Mismatch repair in methylated DNA. Structure and activity of the
  mismatch-specific thymine glycosylase domain of methyl-Cpg-binding
  protein MBD4.";
RT J. Biol. Chem. 278:5285-5291 (2003).
RL J. Biol. Chem. 278:5285-5291 (2003).
CC -1- FUNCTION: Mismatch-specific DNA N-glycosylase involved in DNA
  repair. Has thymine glycosylase activity and is specific for G:T
  mismatches within methylated and unmethylated Cpg sites. Can also
  remove uracil or 5-fluorouracil in G:U mismatches. Has no lyase
  activity. Was first identified as methyl-Cpg-binding protein.
CC -1- SUBUNIT: Interacts with MTH1 (By similarity).
CC -1- SUBCELLULAR LOCATION: Nuclear. In discrete foci.
CC -1- SIMILARITY: Contains 1 methyl-Cpg-binding (MBD) domain.
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CC -----
DR EMBL: AF072249; AAC68878.1; -
DR EMBL: AF120996; AAD56595.1; -
DR EMBL: BC024812; AAH24812.1; -
DR PDB: 1NGN; X-Ray; A=400-554.
DR MGD: MGI:1333850; Mbd4.
DR InterPro: IPR001739; Methyl-Cpg_bind.
DR InterPro: IPR00730; HNH-GPD; 1.
DR Pfam: PF01429; MBD; 1.
DR SMART: SM00391; MBD; 1.
DR PROSITE: PS50982; MBD; 1.
KW 3D-structure; DNA repair; DNA-binding; Hydrolyase; Nuclear protein.
FT DOMAIN 63 135 MBD.
FT DOMAIN 435 498 HNH-GPD.
FT ACT_SITE 534 534 By similarity.
FT CONFLICT 129 129 N -> D (in Ref. 3).
SQ SEQUENCE 554 AA; 62577 MW; 792D37CB180291F5 CRC64;

Query Match 59.6%; Score 1821.5; DB 1; Length 554;
Best Local Similarity 66.2%; Pred. No. 7.6e-103;
Matches 384; Conservative 49; Mismatches 116; Indels 31; Gaps 11;

OY 6 LESLSLGD---RGAPVTYSSERLVPDPNDLRKEDVAMELERVGEDEBOMIKRSSBECN 62
DB 1 MESPNLGDNRVVG-----ESLVPDPDPMDCKEDIDVAGVGVEDGKDLVI--SSERS 50
OY 63 PLLQEPPIASQPGATAGTECRKSVPCGWERVVYKRLFGKTAGRPDVPYFIPISQGLKFRSKS 122
DB 51 SLDGPTAST-LSSTTATGKHKFPVCGWBRVYKRLSGTKACKFPVYFIPISQGLKFRSKR 109
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DB 110 SIANTYLKNGEFTLPBEPFNFVLPRKGSINPKYKQSLAALTSLDPNETDVSQKMLKTRK 169
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DB 170 KKKTDVLPSPGTSPPSSSGISNSNACLRLREHRDIDVDSEKRRKSKRYTVLXGTA 229
OY 243 IKTKKXGCRKSCSGFVQSDSKRESVCKNKADESEPAQSKQLDRVTCISDAQAGETLSV 302
DB 230 SKTKQKCKKSLBSRQRRKRSVYQKVGADRELVQPSQLNRTLCPADACA-RETVGL 288
OY 303 TESENSLVKKKRSLSGSGNFCSEOKTSGTINKFCGSAKSEHNEKEDFLESEELGTGV 362
DB 289 AGE-----EKSPSPGIDLCFIQVTSGTKTKPFSTEAAGANR-EQTFLESEELRSK- 338
OY 363 EYVERK--EHLHTDILKXGSEMDNNSCPFRKDPFTEKI FOEDPTI PRTOIERRKTSLYFSS 420
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Db 339 --GDRKGAHLHTGLVLDGSEMP--SCSQAKKHFTSE--TFQEDSIPTQVEKRKTSLYPSS 394
Qy 421 KYNKEALSPPRRKAFKKWTPRSPFNVLVOETLFFHDPWKLTIATIFLNTSGKMAIPVLMK 480
Db 395 KYNKEALSPPRRKSKFKWTPRSPFNVLVOETLFFHDPWKLTIATIFLNTSGKMAIPVLMK 454
Qy 481 FLEKYPSEAVARTADWRVSELKPLGLYDLRAKTIIVFSDEYLTKQWKYPILHIGIKY 540
Db 455 FLEKYPSEAVARAADWRVSELKPLGLYDLRAKTIIVFSDEYLTKQWKYPILHIGIKY 514
Qy 541 GNDSTYRIFCVNEMKQVHPEDHKLANKYHDMWLNENHEKLSLS 580
Db 515 GNDSTYRIFCVNEMKQVHPEDHKLANKYHDMWLNENHEKLSLS 554

RESULT 3

Q919F1 PRELIMINARY; PRT; 416 AA.
ID Q919F1
AC Q919F1;
DT 01-OCT-2000 (Tremblrel. 15, Created)
DT 01-OCT-2000 (Tremblrel. 15, Last sequence update)
DT 01-OCT-2003 (Tremblrel. 25, Last annotation update)
DE 5-methylcytosine G/T mismatch-specific DNA glycosylase.
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Gallus.
OC NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20512562; PubMed=11058112; DOI=10.1093/nar/28.21.4157;
RA Zhu B., Zheng Y., Anglikar H., Schwarz S., Thiry S., Siegmund M.,
RA Joest J.-P.;
RT "5-Methylcytosine DNA glycosylase activity is also present in the
RT human MBD4 (G/T mismatch glycosylase) and in a related avian
RT sequence."
RL Nucleic Acids Res. 28:4157-4165 (2000).
DR EMBL; AF257107; AAF68981.1; -.
DR HSSP; Q922D7; INGN.
DR InterPro; IPR011257; DNA_glycosylase.
SQ SEQUENCE 416 AA; 45454 MW; A3F70C6FP2133F2A CRC64;

Query Match 28.8%; Score 880.5; DB 2; Length 416;
Best Local Similarity 78.2%; Pred. No. 1.2e-45;
Matches 161; Conservative 20; Mismatches 21; Indels 4; Gaps 2;

Qy 378 RGESEMDNNS--PIRKDTGTEKIFQ--EDTIPRIQIERKTSLYHSSKYNKEALSPPRRK 433
Db 210 RDSADAGGVSWPSDKKSFVAQAPRGTESAPRTQVDRKTKSPYSSKYSKEALSPPRRK 269
Qy 434 AFKKWTPRSPFNVLVOETLFFHDPWKLTIATIFLNTSGKMAIPVLMKFLKYPSEAVART 493
Db 270 AFKKWTPRSPFNVLVOETLFFHDPWKLTIATIFLNTSGKMAIPVLMKFLKYPSEAVART 329
Qy 494 ADWRVSELKPLGLYDLRAKTIIVFSDEYLTKQWKYPILHIGIKYGNDSYRIFCVNEM 553
Db 330 ADMKEMSSILRPLGLYALRAKTIIVFSDEYLTKQWKYPILHIGIKYGNDSYRIFCVNEM 389
Qy 554 KQVHPEDHKLANKYHDMWLNENHEKLSLS 579
Db 390 KQVHPEDHKLANKYHDMWLNENHEKLSLS 415

RESULT 4

Q9SFC1 PRELIMINARY; PRT; 419 AA.
ID Q9SFC1
AC Q9SFC1;
DT 01-MAY-2000 (Tremblrel. 13, Created)
DT 01-MAY-2000 (Tremblrel. 13, Last sequence update)
DT 01-JUN-2003 (Tremblrel. 24, Last annotation update)
DE F17A17.27 protein.
GN Name=F17A17.27;
OS Arabidopsis thaliana (Mouse-ear cress).

OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eustoids II; Brassicales; Brassicaceae; Arabidopsis.
OK NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA Lin X., Kaul S., Town C.D., Benito M.-I., Creasy T.H., Haas B.,
RA Rensing C.M., Kuo H., Fujii C.Y., Utecherack T.R., Barnstead M.E.,
RA Bowman C.L., White O., Niernan W.C., Fraser C.M.;
RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC013463; AAF21203.1; -.
DR HSSP; Q922D7; INGN.
DR GO; GO:0006284; P:base-excision repair; IEA.
DR InterPro; IPR011257; DNA_glycosylase.
DR InterPro; IPR003265; Endo_3c.
DR Pfam; PF00730; Hbh-GPD; 1.
SQ SEQUENCE 419 AA; 47957 MW; BAF0BCA5A710C95 CRC64;

Query Match 11.3%; Score 344; DB 2; Length 419;
Best Local Similarity 25.4%; Pred. No. 6.1e-13;
Matches 99; Conservative 57; Mismatches 146; Indels 88; Gaps 11;

Qy 205 SNFSTHLLKEDGCVDNVFRKVRKKGKVTILKGIPI-----KTKKGGCKKSGSGF 257
Db 88 SNLVSPIADDDDSVSDSHIERQCESEFHEVRRVSPYQGSTVSQSKRGK----- 140
Qy 258 VQSDSKRESVCKNKADES-----EPVAKSQLDRTVCISDAGCGER-----LSVTSE 305
Db 141 ---DS--DSVCSKGGCKRQKQKVRVSPYQASTISQ--CSDLVSSSGGRNKRKSSK 193
Qy 306 ENSLVK-----KERSLSGSGNFCSKQKTSGLINKFCSANDSEHNEKEDTFLSEBEIGT 360
Db 194 RQVVRVRVSPYQGSTVSEQN-----QAPKGLRNYF----- 225
Qy 361 KVEYVERKEHLHTILKRGSEMDNNSCPTRKDPFGKEIPQEDTIPRIQIERKTSLYPSS 420
Db 226 --KVVKSRVYFHADGIVNSESQKESRNVK-----TPVSPVLSLQKTDVYL-- 273
Qy 421 KYNKEALSPPRRKAFKKWTPRSPFNVLVOETLFFHDPWKLTIATIFLNTSGKMAIPVLMK 480
Db 274 -----RKTDNTWVTPRSPFNVLVOETLFFHDPWKLTIATIFLNTSGKMAIPVLMK 323
Qy 481 FLEKYPSEAVARTADWRVSELKPLGLYDLRAKTIIVFSDEYLTKQWKYPILHIGIKY 540
Db 324 LFGCTDAKATATEVKEEIEILKPLGLQKKRTMIGRLSLEYQBSWHTVQLHGVGKY 383
Qy 541 GNDSTYRIFCVNEMKQVHPEDHKLANKYHDM 570
Db 384 AADAYALFCNGMWDVRVKNHMLNLYWDYL 413

RESULT 5

Q84WT3 PRELIMINARY; PRT; 407 AA.
ID Q84WT3
AC Q84WT3;
DT 01-JUN-2003 (Tremblrel. 24, Created)
DT 01-JUN-2003 (Tremblrel. 24, Last sequence update)
DT 01-OCT-2003 (Tremblrel. 25, Last annotation update)
DE Hypothetical protein At3g07930.
GN Name=At3g07930;
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eustoids II; Brassicales; Brassicaceae; Arabidopsis.
OK NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA Yamada K., Chan M.M., Chang C.H., Dale J.M., Hsuan V.W., Lee J.M.,
RA Onodera C.S., Quach H.L., Tang C., Toriumi M., Wong C., Wu H.C.,
RA Yu G., Yuan S., Carninci P., Chen H., Cheuk R., Hayashizaki Y.,
RA Ishida J., Jones T., Kamiya A., Kawai J., Kim C.J., Narusaka M.,
RA Nguyen M., Palm C.D., Sakurai T., Satou M., Seki M., Shinu P.,
RA Southwick A., Tripp M.G., Wu T., Shinozaki K., Davis R.W., Ecker J.R.,

RA Theologis A.;
 RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL: BT002799; AA022623.1; -.
 DR HSBP; Q922D7; INGN.
 DR GO: GO:0006284; P:base-excision repair; IEA.
 DR InterPro: IPR011257; DNA_glycosylase.
 DR InterPro: IPR003465; Endo_3c.
 DR Pfam: PF00730; HNH-GPD; 1.
 DR Hypothetical protein.
 KM
 SQ SEQUENCE 407 AA; 46941 MW; DD758CD862EF54F CRC64;

Query Match 11.1%; Score 338; DB 2; Length 407;
 Best Local Similarity 26.6%; Pred. No. 1,4e-12;
 Matches 102; Conservative 51; Mismatches 155; Indels 76; Gaps 11;

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QY 251 RKSQSGFVQ-SDSKRESVGNKADASEPVAQKSQLDRVCISDAGACGELLSTSEENSL 309
DB 30 RRPDDDFIEVDENSNFALFKEDDEK-----NRDLGLVDDGSTNLVLQCHDDGSL 80
QY 310 YKKKERSLS---SGSNFCSEQR-----TSGIINFKCSAKDSEHNEKEDTFLESE 356
DB 81 EKDNSSLSDDLPSGFYKGVKRRKRDDFGSITTSNLSVQIADDD---DDSVSDSHIERO 137
QY 357 EIGTKVEYVERK-----EHLHTDLKRGSEMDN-NCSPTRKDTGTEKI---FOEDT 403
DB 138 ECSKVQAKVPRVSPYFOASTISQCDSDIVSSQSGRNKRGSSKQVKARARSPYFOEST 197
QY 404 IP-----RTQIERKRTSLVF-----SSKYNKEALSP----- 430
DB 198 VEEQNPQAFKGLNRYFKVYKSRFYADGIQVNESQKESRVRKTPPIYSPULSOKTD 257
QY 431 ---RRKAFKKWTTPRSFNNLVQETLFHDPMKLLATIFLNRTSGKMAIPVLMKELEKP 486
DB 258 DYVLRKTPNTWVPPSPSCNLLQEDHMDPMRVLVICMLNTKTSGAQTGVSIDLFGLCT 317
QY 487 SAEVATADWRDVSSELKPLGLYDLRAKTIYKPSDYLTKOMKPIYELHGLICRGNDYSR 546
DB 318 DAKTATEVEKEEIEENLIKPLGLQKKRTKMIORLSLEYLOESWTHVLQHGKADADAYA 377
QY 547 IFCVNEWKQVHPEDHLNKYHML 570
DB 378 IFCGNMWDRKPNHMLNTYMDYL 401

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RESULT 6

Q7T2T7 PRELIMINARY; PRT; 524 AA.

AC 07T2T7;
 DT 01-OCT-2003 (TREMBLrel. 25, Created)
 DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
 DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
 DE Methyl-cytosine binding protein 2.
 GN Name=Mecp2;
 OS Brachydanio rerio (Zebrafish) (Danio rerio).
 OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Danio.
 OC NCB1_TaxID=7955;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Coverdale L.E., Martin C.C.;
 RL Submitted (OCT-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AY298900; AAP57248.2; -.
 DR HSSP; Q9UN29; 11G4.
 DR ZFIN: ZDB-GENE-030131-7199; mec2.
 DR GO: GO:0003677; F:DNA binding; IEA.
 DR InterPro: IPR001739; Methyl-Cpg_bind.
 DR Pfam: PF01429; MBD; 1.
 DR SMART; SM00391; MBD; 1.
 SQ SEQUENCE 524 AA; 57152 MW; B8593B4B84DCD21 CRC64;

Query Match 7.8%; Score 238.5; DB 2; Length 524;
 Best Local Similarity 24.4%; Pred. No. 2.2e-06;

Matches 106; Conservative 57; Mismatches 159; Indels 113; Gaps 18;

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QY 27 VPDPNDLRKDDVAMELERVEDEEQMMIKRSSCNPLQBPISAOFGA---TAGTECR 83
DB 49 VPPPSLEFTQDVQQAQ-AGKSE-----PI--DPEVGAISAESSASAKOR 93
QY 84 KSV-----PCGWERVVKORLFGTAGRFVYFISPOLKFRSKSLANYLAK 130
DB 94 RSVIRDRGPMTEDSLPGWTRKIKQKSKSGSACKFDVYLINPGKAFRSKVELMAYRQK 153
QY 131 NGESTLKEDEPDPFVLSKRGIKSRVYKDCSMALTSHLQNGSNNSNMLRTSKCKQVFM 190
DB 154 VGDITTDNDPDPFVY-TGRGSPSR-----RRKRPKKPKRV 188
QY 191 PPSSESSELRGSLNFTSHLLKEDBGVDVNPFRVKKPKGKVTI----- 237
DB 189 KPS-----GRGRPRKSGKVRQATEGV--AVKRVLEKPGKLVMPVAPPTERGA 239
QY 238 -LKGIPIKTKKKGKSGFVQSDSKRESVGNKADASEPVAQKSQLDRVCISDAGAC 296
DB 240 PLGQAPVAKARGRK-----KSEQDPSTPKRGKRPATVSQS---TVGTSAAY 288
QY 297 GETLSVTSE-----ENSLVKKERSLSGSGNFCEQKTSGLINKFCSAKSEHNEKYE 349
DB 289 AAAAILTFAKKKALKESSAKPVQGRALP-----IKKRTRETLIEL-EASTTSATETPE 342
QY 350 DTFLESEIEIGTKVEY-VERKEHLHTDLKRGSEMDNNSCPRKDTGTEKIQEDTIPRTQ 408
DB 343 KLTASTVTPTGESELETQCKPHKPS--RKIKHEADPGSSSGSTTASG-----VAPKSH 393
QY 409 IERRKTSLYFSKYN 423
DB 394 KKRDRGQHPFKHHH 408

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RESULT 7

Q9YGC6 PRELIMINARY; PRT; 467 AA.

AC Q9YGC6;
 DT 01-MAY-1999 (TREMBLrel. 10, Created)
 DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)
 DT 05-JUL-2004 (TREMBLrel. 27, Last annotation update)
 DE Methyl-Cpg-binding protein Mecp2 (Methyl-Cpg-binding protein 2).
 GN Name=Mecp2;
 OS Xenopus laevis (African clawed frog).
 OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
 OC Xenopodinae; Xenopus.
 OC NCB1_TaxID=8355;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=98282101; PubMed=9620779; DOI=10.1038/561;
 RX Jones P.L., Veenstra G.J.C., Wade P.A., Vermaak D., Kass S.U.,
 RA Landsberger N., Strouboulis J., Wolffe A.P.;
 RT "Methylated DNA and Mecp2 recruit histone deacetylase to repress
 transcription.";
 RT Nat. Genet. 19:187-191 (1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Kass S.U., Strouboulis J., Wolffe A.P.;
 RL Submitted (FEB-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF106951; AAD03736.1; -.
 DR EMBL: AF051768; AAD02651.1; -.
 DR HSSP; P51608; 10X9.
 DR GO: GO:0005634; C:nucleus; IEA.
 DR GO: GO:0003677; F:DNA binding; IEA.
 DR GO: GO:0006355; P:regulation of transcription; IEA.
 DR InterPro: IPR00637; A+T hook.
 DR InterPro: IPR001739; Methyl-Cpg_bind.
 DR Pfam: PF02178; AT_hook; 1.
 DR Pfam: PF01429; MBD; 1.
 DR SMART; SM00384; AT hook; 2.
 DR SMART; SM00391; MBD; 1.

SQ SEQUENCE 467 AA; 51757 MW; 5D3A719A59E560BC CRC64;
Query Match 7.2%; Score 220.5; DB 2; Length 467;
Best Local Similarity 25.7%; Pred. No. 2.3e-05;
Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
QY 22 SSERLVPPNDLRKEDVAMELER---VGEDEEQWIKRS--SECNPLOPIASAOQGA 76
DB 49 SSEH-QPEPADDEGADMSBAEENLVPESSASAKGRSVIRDGPRYEDP----- 99
QY 77 TAGTECRKSVPCGMEVRYVQRIPEKTAGRPDVYFISPOGLKPKRSKSLANYLHKNETSL 136
DB 100 -----TLPEGMWTRKLRKGRSGRSACKFPVYLLINPGKAFRSKVELIAVFOKGDFTSL 151
QY 137 KPREDFTVLKSRGKSKRYXOCSMALTSHLQONSNMNLFRPSKKOKOV--FMPSS 194
DB 152 DPNPDFTV-TGRGSPSRREQ-----KQPKPKPKPKSSVSGRGGRPKGSIKKVPPVK 204
QY 195 SSRLQESHRLNFTSTHLTK----EDEGVDDVAFKRV-----RKPQKVTILKGI 242
DB 205 SEGQVVKVIERK-SFGKLVKMPYSGTKEASDATTSCQVLYIKGGRKRGSE-IDPSAP 262
QY 243 IKTTKGCRKSCSGFVQSDSRSEVYCNADABSEFVQAKSOLDRVCISDAGCGETLSV 302
DB 263 KKGKGRKPSNVSLAAMAAAKKAI---KESSIKPLE-----TVLPTRKRTRETTISV 313
QY 303 TSEE-----NSLVKK-----KERSLSGSGNFCSEBKTSIGIINKPCASDSEHNEKYE 349
DB 314 DVKQTIKEPLTPVLEKWMKQNPSPSRSTEGSPKTKLGPCKELQHHHHHHHHH 373
QY 350 DTFLSEERIGTKVEVERKEHL 371
DB 374 HHSESKASATSPPEPSTKONI 395
RESULT 8
Q7SCQ2 PRELIMINARY; PRT; 682 AA.
AC Q7SCQ2:
DT 01-MAR-2004 (TREMBlrel. 26, Created)
DT 01-MAR-2004 (TREMBlrel. 26, Last sequence update)
DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
DE Hypothetical protein.
GN Name=NCU09815.1;
OS Neurospora crassa.
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
OC Sordariomycetidae; Sordariales; Sordariaceae; Neurospora.
OX NCBI_TaxID=5141;
RN RP
RP SEQUENCE FROM N.A.
RC STRAIN=OR74A;
RA Galagan J.E., Calvo S.E., Borkovich K.A., Selker E.U., Read N.D.,
RA Jaffe D., Fitzhugh W., Ma L.-D., Smirnov S., Purcell S., Reiman B.,
RA Elkins T., Engels R., Wang S., Nielsen C.B., Butler J., Endrizzi M.,
RA Qui D., Ianakiev P., Pedersen D., Nelson M., Washburne M.,
RA Selitrenikoff C.P., Kinsey J.A., Braun E.L., Zelter A., Schulte U.,
RA Koche G.O., Jedd G., Mewes W., Staben C., Marcotte E., Greenberg D.,
RA Roy A., Foley K., Naylor J., Thomann N., Barrett K., Gnetre S.,
RA Kamal M., Kamyseilis M., Mancelli E., Bielke C., Rudd S., Frisman D.,
RA Krysstofova S., Rasmussen C., Metzberg R.L., Perkins D.D., Kroken S.,
RA Cogoni C., Macino G., Catchside D., Li W., Pratt R.J., Osmari S.A.,
RA Desouza C.C., Glass L., Orbach M.J., Berglund J., Voelker R.,
RA Yarden O., Plamann M., Seiler S., Dunlap J., Radford A., Aramayo R.,
RA Natvig D.O., Alex L.A., Mannheim G., Eboile D.J., Freitag M.,
RA Paulsen I., Sachs M.S., Lander E.S., Nusbaum C., Birren B.,
RA "The Genome Sequence of the Filamentous Fungus Neurospora crassa."
RT Nature 0:0-0(2003).
CC -!- CAUTION: The sequence shown here is derived from an
EMBL/GenBank/DBJ whole genome shotgun (WGS) entry which is
preliminary data.
CC EMBL; AABX01000087; EAA34519.1; -
DR GO; GO:0006284; P:base-excision repair; IEA.
DR InterPro; IPR011257; DNA_glycosylase.

DR InterPro; IPR003265; Endo_3c.
DR Pfam; PF00730; Hnh-GPD; 1.
KW Hypothetical protein.
SQ SEQUENCE 662 AA; 75928 MW; 4FDCCAA26102EB84 CRC64;
Query Match 7.0%; Score 213; DB 2; Length 682;
Best Local Similarity 20.8%; Pred. No. 0.0011;
Matches 93; Conservative 62; Mismatches 155; Indels 138; Gaps 14;
QY 240 GIPIKTKGCRKSCSGF-VQSDSKRS-----VNKADABSEFPAQKSQ 284
DB 164 GKRVKTKTGWGGRVSRFMAQDDPAKTSQVTRGRTGVSGAVRKAMNGHEPISQGGK- 222
QY 285 DRVCISDAGAGCBTTLVTSEENSLVKKK-RSLSSGNSNFCSEBKTSIGIINKPCASDSE 343
DB 223 -----MLPTRWMSVVRKSAGPSGGBE---SKAKSRLLDK-TKLKHSK 263
QY 344 HNEK-YEDTFLESEERIGTKVEVERKEHLHTDLKSGSEMUNNC-SPTKDFTEKIFQ 400
DB 264 FPDEPSYADTWSPQSLASHSSPNEMLDVALGDVSVAGTSSPDECVLAANVRITGERITYK 323
QY 401 EDTI---PRTQIERKTSLYFSSKYNKEALSPRRKAFKKTPTP-----RSPFN 446
DB 324 SPFSEBPPMPPISTSKASPPKSTKTSKSSPTKKS--RNHPRGISCLPIAPISARFG 381
QY 447 LVQETLPHDPWKLILATIFLNRTSGKMAIPVLMKFLKYSAAEVARTADWRDVSSELKPL 506
DB 382 LIQEVAAADPRLILATVFLIKRGMTAIPFLRQMDLFTPEALASADSEIINLRPL 441
QY 507 GLYDLRAKTIKVSF-----DEYLTQWKYPIELHGI----- 537
DB 442 GLSVNRCSVLQKVARMFIECPCKEKRYGVKNYPRPDAGVAGKVGQFTGEPDFHIGKA 501
QY 538 -----GKYGNSYRIFCN----- 551
DB 502 SQAEFDDDDDRINAIRKAKERHAIGCAWEIGLTLQGPALDSWRIFCRDKLLGRADWKGK 561
QY 552 -----EMKQVPEBDKINKYHMDLM 571
DB 562 GRHPEQPEMWRVLPQDKELRAYLRMMW 589
RESULT 9
MEC2 HUMAN
ID MEC2_HUMAN STANDARD; PRT; 486 AA.
AC P51608; O15233;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-CpG-binding protein 2 (MeCP-2 protein) (MeCP2).
GN Name=MECP2;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxID=9606;
RN RP
RP SEQUENCE FROM N.A.
RA Kudo S., Fukuda M.,
RL Submitted (SEP-1995) to the EMBL/GenBank/DBJ databases.
RN RP
RP SEQUENCE FROM N.A.
RC Tissue=Placenta;
RA Thiesen J., Straetling W.H.,
RL Submitted (APR-1997) to the EMBL/GenBank/DBJ databases.
RN RP
RP SEQUENCE FROM N.A.
RX MEDLINE=97130625; PubMed=9976388;
RA Vilain A., Apioy F., Vogt N., Dutrillaux B., Malfoy B.,
RT "Assignment of the gene for methyl-CpG-binding protein 2 (MECP2) to
RL human chromosome band Xq28 by in situ hybridization."
RN CytoGenet. Cell Genet. 74:293-294(1996).
RP SEQUENCE FROM N.A.

RA Reichwald K., Rosenthal A., Kioschis P., Platzter M.,
 RT "Mapping and sequence analysis of the human MECP2 locus.",
 RL Submitted (Oct-1997) to the EMBL/GenBank/DBD databases.
 RN [15]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=9929240; PubMed=10369871; DOI=10.1093/hmg/8.7.1253;
 RA Coy J.F., Sedlacek Z., Baechner D., Delius H., Poustka A.,
 RT "A complex pattern of evolutionary conservation and alternative
 RT polyadenylation within the long 3'-untranslated region of the methyl-
 RT Cpg-binding protein 2 gene (MECP2) suggests a regulatory role in gene
 RT expression.";
 RL Hum. Mol. Genet. 8:1253-1262 (1999).
 RN [16]
 RP SEQUENCE FROM N.A.
 RA Reichwald K., Thiesen J., Wiehe T., Weitzel J., Straetling W.H.,
 RA Kioschis P., Rosenthal A., Platzter M.,
 RT "Comparative sequence analysis of the MECP2-locus in human and mouse
 RT reveals new untranslated regions".
 RL Submitted (JUN-1999) to the EMBL/GenBank/DBD databases.
 RN [17]
 RP SEQUENCE FROM N.A.
 RC TISSUE=placenta;
 RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Straubeberg R.L., Reingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diachenko L., Marsina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stalonen M., Soares M.B., Bonaldi M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Uedtin T.B., Toshlyuk S., Carninci P., Prange C.,
 RA Rana S.S., Loquellano N.A., Peters G.J., Abtenson R.D., Mallary S.J.,
 RA Bobak S.A., Mewman P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Morley K.C., Hale S., Garcia A.M., Gay L.J., Huiyk S.W.,
 RA Villalón D.K., Muray D.M., Sodergren B.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Ketterman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.W.,
 RA Butcherfield Y.S.N., Krzywinski M.I., Skalska U., Smalins D.E.,
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.,
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 RN [18]
 RP SEQUENCE OF 10-486 FROM N.A.
 RC TISSUE=skeletal muscle;
 RX MEDLINE=96327611; PubMed=8672133;
 RA D'Esposito M., Quadert N.A., Ciccodicola A., Bruni P., Esposito T.,
 RA D'Urbio M., Brown S.D.M.,
 RT "Isolation, physical mapping, and Northern analysis of the X-linked
 RT human gene encoding methyl Cpg-binding protein, MECP2.";
 RL Mamm. Genome 7:533-535 (1996).
 RN [19]
 RP SEQUENCE OF 10-486 FROM N.A.
 RA Reichwald K., Bauer D., Brenner V., Drescher B., Coy J.F.,
 RA Kioschis P., Korn B., Myakatura G., Platzter M., Poustka A.,
 RA Sandoval N., Rosenthal A.,
 RT Submitted (DEC-1996) to the EMBL/GenBank/DBD databases.
 RN [10]
 RP REVIEW ON VARIANTS.
 RX PubMed=12872250; DOI=10.1002/humu.10243;
 RA Mittenberger-Miltenyi G., Laccione F.,
 RT "Mutations and polymorphisms in the human methyl Cpg-binding protein
 RT MECP2.";
 RL Hum. Mutat. 22:107-115 (2003).
 RN [11]
 RP VARIANTS RTT TRP-106, CYS-133, SER-155, MET-158 AND CYS-306, AND
 RP VARIANT LYS-397.
 RX PubMed=10577905;
 RA Amir R.E., Budden S., Naidu S., Pereira J.L.P., Lo I.F.M.,
 RA Zoghbi H.Y., Schanen N.C., Francke U.,
 RT "Ret syndrome and beyond: recurrent spontaneous and familial MECP2
 RT mutations at Cpg hotspots.";
 RL Am. J. Hum. Genet. 65:1520-1529 (1999).
 RN [12]
 RP VARIANTS RTT TRP-106, CYS-133, SER-155 AND MET-158.
 RX MEDLINE=99438392; PubMed=10508514; DOI=10.1038/13810;
 RA Amir R.E., Van den Veyver I.B., Wan M., Tran C.Q., Francke U.,
 RA Zoghbi H.Y.,
 RT "Ret syndrome is caused by mutations in X-linked MECP2, encoding
 RT methyl-Cpg-binding protein 2.";
 RL Nat. Genet. 23:185-188 (1999).
 RN [13]
 RP INVOLVEMENT IN X-LINKED MENTAL RETARDATION WITH PROGRESSIVE
 RP SPASTICITY.
 RX PubMed=1036043;
 RA Meloni I., Brucini M., Longo I., Mari F., Rizzolio F., D'Adamo P.,
 RA Denvirand K., Fryns J.-P., Tonello D., Renieri A.,
 RT "A mutation in the Ret syndrome gene, MECP2, causes X-linked mental
 RT retardation and progressive spasticity in males.";
 RL Am. J. Hum. Genet. 67:982-985 (2000).
 RN [14]
 RP VARIANTS RTT VAL-100, GLN-106, TRP-106, CYS-133, ARG-152, SER-155;
 RP MET-158; ARG-305; CYS-306 AND HIS-306, AND VARIANTS CYS-86; MET-203;
 RP PRO-287; ALA-291; LYS-397; ILE-412 AND THR-444.
 RX PubMed=11055898;
 RA Buysse I.M., Pang P., Hoon K.T., Amir R.E., Zoghbi H.Y., Roa B.B.,
 RT "Diagnostic testing for Ret syndrome by DHPLC and direct sequencing
 RT analysis of the MECP2 gene: identification of several novel mutations
 RT and polymorphisms.";
 RL Am. J. Hum. Genet. 67:1428-1436 (2000).
 RN [15]
 RP VARIANT MEX16 VAL-140, AND VARIANT MET-203.
 RX MEDLINE=20465115; PubMed=11007980; DOI=10.1016/S0014-5793(00)01994-3;
 RA Oriccio A., Lam C., Galli L., Dotti M.T., Hayek G., Tong S.F.,
 RA Poon P.M., Zappella M., Federico A., Sorrentino V.,
 RT "MECP2 mutation in male patients with non-specific X-linked mental
 RT retardation.";
 RL FEBS Lett. 481:285-288 (2000).
 RN [16]
 RP VARIANTS RTT LEU-101; HIS-101; THR-101; TRP-106; CYS-133; CYS-134;
 RP ARG-152; MET-158; ARG-225; LEU-302; CYS-306 AND HIS-306, AND VARIANTS
 RP LEU-229 AND THR-439.
 RX PubMed=10767337; DOI=10.1093/hmg/9.7.1119;
 RA Chesdale J.P., Gill H., Fleming N., Maynard J., Kerr A., Leonard H.,
 RA Kravczak M., Cooper D.N., Lynch S., Thomas N., Hughes H., Hulten M.,
 RA Ravine D., Sampson J.R., Clarke A.,
 RT "Long-read sequence analysis of the MECP2 gene in Ret syndrome
 RT patients: correlation of disease severity with mutation type and
 RT location.";
 RL Hum. Mol. Genet. 9:1119-1129 (2000).
 RN [17]
 RP VARIANTS RTT GLN-106; MET-158; ARG-302; CYS-306 AND ALA-322.
 RX PubMed=10814719; DOI=10.1093/hmg/9.9.1377;
 RA Bienvenu T., Carrie A., de Roux N., Vinet M.-C., Jonveaux P.,
 RA Couvert P., Villard L., Arizmanoglou A., Beldjord C., Fontes M.,
 RA Tardieu M., Chelly J.,
 RT "MECP2 mutations account for most cases of typical forms of Ret
 RT syndrome.";
 RL Hum. Mol. Genet. 9:1377-1384 (2000).
 RN [18]
 RP VARIANTS RTT MET-158; HIS-302 AND CYS-306, AND VARIANTS VAL-201;
 RP ALA-232; LEU-251 AND SER-376.
 RX PubMed=10944854;
 RA Amano K., Nomura Y., Segawa M., Yamakawa K.,
 RT "Mutational analysis of the MECP2 gene in Japanese patients with Ret
 RT syndrome.";
 RL J. Hum. Genet. 45:231-236 (2000).
 RN [19]
 RP VARIANTS RTT TRP-106, PHE-124, CYS-133, CYS-134, ARG-152, MET-158 AND
 RP CYS-306.
 RX MEDLINE=20439334; PubMed=10991688;
 RA Obata K., Matsui T., Yamashita Y., Fukuda T., Kuwajima K.,
 RA Horiuchi I., Nagamitsu S., Iwanaga R., Kimura A., Omori I., Endo S.,
 RA Mori K., Kondo I.,

Query Match	6.8%	Score 208.5	DB 2	Length 498
Beet Local Similarity	23.9%	Pred. No. 0.00014		
Matches 104	Conservative 53	Mismatches 149	Indels 129	Gaps 17
QY	37	EDVAMELEIRVGDEDEKQWMIKRSSECNPLLOEPISAOFG	-----ATAGT	80
DB	34	KDKPLKPKKVKKKDKKEKEGKEHPEYQPSAHHSABEAKGKATSESGSAPAVPEASASP	93	
QY	81	ECKRSV-----PGMERVYKQRLFGKTAGRFVDYFISPOGLKFRSKSLANY	127	
DB	94	KQRSIIIRDRGPMDDPTLPFGMTKRLKQKRSRGAGKYDVLINPOGAFRSKVETLAV	153	
QY	128	LHKNGESTLKEDPDFTLTKRGKISRKKDCSMALTSHLQNSNNSMNLTRSKCKKD	187	
DB	154	PEKVGDTLDPDDEPFTV-TGSGSPSR-----REQRPKK-	187	
QY	188	VFMPPSSSSSEIQESRGL---SNFTSTHLLKXEDGVDDVNFKKVKKPKGKVTILKGIPIK	244	
DB	188	-----PKSKAKGTGTGRGRPKSGTTRKKAATSEVQVK--RVLEKSPGK--LVKMP-	238	
QY	245	KTKKGCRKSCSGFVQSDS---KRESVCNKDAESEPVAAQKSQLDRTVCISDAGAGETL	300	
DB	229	QTSFGKAEKGAGATSTQVMVYKRPGRKKAADPOAIPKK-----GKRPGSV	288	
QY	301	SVTSESNLVKKKREKSLSSGNSFCGEQKTSGLINFCGAKDSHNEKXEDFTLSEBEIGT	360	
DB	289	AAAAAEAKKAVKESIR-----SVQETVLPIKK-----RKRET-----V	324	
QY	361	KVEVVERKEHLHTDL--KRGSEMNNKSPFKDPTGKIFQEDTIPRTQIERKTSLYF	418	
DB	325	SIEVEAVKPLIVLSTLGEKSGKGLTKCSPGK-----SKESPICK-----	366	
QY	419	SSKYNKEALSPPRK 433		
DB	367	-----SSASSPPKKE 377		
RESULT 11				
Q72384				
ID	PRELIMINARY	PRT	516 AA.	
AC	Q72384			
DT	01-OCT-2003 (TREMBLrel. 25, Created)			
DT	01-OCT-2003 (TREMBLrel. 25, Last sequence update)			
DT	01-MAR-2004 (TREMBLrel. 26, Last annotation update)			
DE	Hypothetical protein DKFZp686A24160 (Fragment).			
GN	Name=DKFZp686A24160;			
OS	Homo sapiens (Human)			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.			
OX	NCBI_TaxID=9606;			


```

[1]
RN SEQUENCE FROM N.A.
RC TISSUE=human colon endothelial primary cell culture;
RA Bloeker H., Boecker M., Mewes H.W., Well B., Amid C., Osanger A.,
RA Robo G., Han M., Wiemann S.;
RA Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BX538060; CAD97991.1; -.
DR HSP; Q9UN29; 1164.
DR GO; GO:0005634; C:nucleus; IEA.
DR GO; GO:0003677; F:DNA binding; IEA.
DR GO; GO:0003555; P:regulation of transcription, DNA-dependent; IEA.
DR InterPro; IPR000637; A+T hook.
DR InterPro; IPR001739; Methyl-CpG_bind.
DR Pfam; PF02178; AT hook; 1.
DR Pfam; PF01429; MBD; 1.
DR SMART; SM00384; AT hook; 2.
DR SMART; SM00391; MBD; 1.
KW Hypothetical protein.
FT NON_TER
SQ SEQUENCE 516 AA; 55204 MW; 27CD37B9164176B0 CRC64;

Query Match
Best Local Similarity 23.9%; Score 208.5; DB 2; Length 516;
Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;

QY 37 EDVAMELERVGEDEBQMIKRSSECNPLLOEPASQFG-----ATAGT 80
DB 52 KKKPLKFKVKKDKKEKKGKHEPVQPSAHNSAEPABAGKAESESGSAPAVPEASAP 111
QY 81 ECRKSV-----PCGWERVVKQRLFGKTAGRDVYFISQGLKFRSKSLANY 127
DB 112 KQRRSIIIRDRGPMYDPTLPBEGWTRKLKQKRSGRAGKDYVLLINQGAFRSKVELLAY 171
QY 128 LKNGETSLKPEDPFTVLSKRGIRKYDCSMALTSHLQNSNNMNLRTSKCKD 187
DB 172 FEKVGDTSLDNDPFTV-TGRGSPR-----REQRPKK- 205
QY 188 VFMPPSSSELOESRGL---SNFTSTHLLKDEGVDDVNFRRKVRKPKGKVTILKGIPIK 244
DB 206 ----PKSPKAPGTGRGRPKSGGTRPKAATSEGVQK--RVLEKSPGK--LLVMMP- 256
QY 245 KTKKGRKSCSGFVQDS---KRSVCNKADABEPVAKQSLDRTVCISDAGAGETL 300
DB 257 QTSFGKAGGAGGATTTQVMVTKRPGKRAEADPQAIPIKGR-----GRKGSYV 306
QY 301 SVTSEENSLVKKKERSLSSGNSFCSEOKTSGIINKFCSAKDSHNEKVEDTFLSEIEGT 360
DB 307 AAAAAAAKKAVKESIR-----SVQETVLPKK-----RKRET-----V 342
QY 361 KVEVERKEHLHTDIL--KRGSEMDNNSCFTRKDFTEGKIFQEDTTPRTQIERRTSLYF 418
DB 343 SIEVEKVVPLVSTLGEKSGKGLTKCKSPGRK-----SKESPCKGR----- 384
QY 419 SKRTKEALSPRRK 433
DB 385 ----SSSASSPPKCE 395

RESULT 12
MEC2_MACEFA STANDARD; PRT; 486 AA.
AC 0951G8;
DT 25-OCT-2004 (Rel. 45, Created)
DT 25-OCT-2004 (Rel. 45, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-CpG-binding protein 2 (Mecp-2 protein) (Mecp2).
GN Name=Mecp2;
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecinae; Macaca.
OC NCBI_taxid=9541;
RN [1]

```

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RP SEQUENCE FROM N.A.
RA Muramatsu S.;
RT "Excessive hand-wringing in a MTPP-created monkey.";
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
CC -! FUNCTION: Chromosomal protein that binds to methylated DNA. It can
CC bind specifically to a single methyl-CpG pair. It is not
CC influenced by sequences flanking the methyl-CpGs. Mediates
CC transcriptional repression through interaction with histone
CC deacetylase and the corepressor SIN3A.
CC -! SUBUNIT: Interacts with FMBP3 (By similarity).
CC -! SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-CpG in the
CC genome (By similarity).
CC -! SIMILARITY: Contains 2 A+T hook DNA-binding repeats.
CC -! SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; AF295597; AAK97131.1; -.
DR HSP; P51608; 10K9.
DR InterPro; IPR000637; AT hook.
DR InterPro; IPR001739; Methyl-CpG_bind.
DR Pfam; PF02178; AT hook; 1.
DR Pfam; PF01429; MBD; 1.
DR SMART; SM00384; AT hook; 2.
DR SMART; SM00391; MBD; 1.
DR PROSITE; PS50982; MBD; 1.
KW DNA-binding; Nucleic protein; Repeat; Repressor;
KW Transcription regulation.
FT DOMAIN 90 162 MBD.
FT DNA_BIND 185 197 A+T hook 1.
FT DNA_BIND 265 277 A+T hook 2.
FT DOMAIN 366 372 His-rich.
FT DOMAIN 376 405 Pro-rich.
SQ SEQUENCE 486 AA; 52426 MW; 3471B61D90D92A7D CRC64;

Query Match
Best Local Similarity 23.9%; Score 207.5; DB 1; Length 486;
Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;

QY 37 EDVAMELERVGEDEBQMIKRSSECNPLLOEPASQFG-----ATAGT 80
DB 22 KKKPLKFKVKKDKKEKKGKHEPVQPSAHNSAEPABAGKAESESGSAPAVPEASAP 81
QY 81 ECRKSV-----PCGWERVVKQRLFGKTAGRDVYFISQGLKFRSKSLANY 127
DB 82 KQRRSIIIRDRGPMYDPTLPBEGWTRKLKQKRSGRAGKDYVLLINQGAFRSKVELLAY 141
QY 128 LKNGETSLKPEDPFTVLSKRGIRKYDCSMALTSHLQNSNNMNLRTSKCKD 187
DB 142 FEKVGDTSLDNDPFTV-TGRGSPR-----REQRPKK- 175
QY 188 VFMPPSSSELOESRGL---SNFTSTHLLKDEGVDDVNFRRKVRKPKGKVTILKGIPIK 244
DB 176 ----PKSPKAPGTGRGRPKSGGTRPKAATSEGVQK--RVLEKSPGK--LLVMMP- 226
QY 245 KTKKGRKSCSGFVQDS---KRSVCNKADABEPVAKQSLDRTVCISDAGAGETL 300
DB 227 QTSFGKAGGAGGATTTQVMVTKRPGKRAEADPQAIPIKGR-----GRKGSYV 276
QY 301 SVTSEENSLVKKKERSLSSGNSFCSEOKTSGIINKFCSAKDSHNEKVEDTFLSEIEGT 360
DB 277 AAAAAAAKKAVKESIR-----SVQETVLPKK-----RKRET-----V 312
QY 361 KVEVERKEHLHTDIL--KRGSEMDNNSCFTRKDFTEGKIFQEDTTPRTQIERRTSLYF 418
DB 343 SIEVEKVVPLVSTLGEKSGKGLTKCKSPGRK-----SKESPCKGR----- 354

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QY 419 SSKYNKEALSPRRK 433
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Db 355 -----SSASPPKKE 365

RESULT 13
042403 PRELIMINARY; PRT; 344 AA.
ID 042403
AC 042403;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Attachment region binding protein (Fragment).
GN Name=ARBP;
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Gallus.
OX NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97415642; PubMed=9271441;
RA Weitzel J.M., Burmeister H., Straetling W.H.;
RT "Chicken MAR-binding protein ARBP is homologous to rat methyl-CpG
RL Mol. Cell. Biol. 17:5656-5666(1997).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=20188769; PubMed=10723722;
RA Reichwald K., Theissen U., Wiene T., Weitzel J., Straetling W.H.,
RA Kioschis P., Pousetka A., Rosenthal A., Platzer M.;
RT "Comparative sequence analysis of the MEC22-locus in human and mouse
RL reveals new transcribed regions.";
Mamm. Genome 11:182-190(2000).
DR EMBL; Y14166; CAA74577.1; -.
DR PDB; 10B1; NMR; A=64-196.
DR GO; GO:0005634; C:nucleus; IEA.
DR GO; GO:0003700; F:transcription factor activity; IEA.
DR GO; GO:0006355; F:regulation of transcription, DNA-dependent; IEA.
DR InterPro; IPR006337; A+T hook.
DR InterPro; IPR001739; Methyl-CpG_bind.
DR InterPro; IPR005976; Wilms_tumour.
DR Pfam; PF01429; MBD; 1.
DR PRINTS; PR00929; ATHOOK.
DR PRINTS; PR00049; WILMSTMOUR.
DR SMART; SM00391; MBD; 1.
FT NON TER 344
SQ SEQUENCE 344 AA; 33640 MW; 53DD7BD9C9DF4FE3 CRC64;

Query Match 6.6%; Score 202.5; DB 2; Length 344;
Best Local Similarity 34.7%; Pred. No. 0.0002;
Matches 59; Conservative 17; Mismatches 57; Indels 37; Gaps 5;

QY 16 AAPVTTSERLVPPENDLRKEDVAMELER-----VGDEBQMIKRSSECNPLIQEPIA 70
   : |||:::
Db 5 AAAAAGGHERL-----EQADGEGVAGLKERPPKAKGKRERDEAEAEAPGAPAE 59

QY 71 SAQGATGTGTEC-----RKSV-----PCGMEVVKORLFGKTAG 104
   : : : : :
Db 60 AGKADGSGGTAAAPAVPEASAPKQRRSIIRDRGPMYDPTLPBGWTRKLRKQKSGRSAG 119

QY 105 RFDVVFIPQGLKFRKSSILANYLHNGETSLKPEDPFTVLSKRGKISR 154
   : : : : :
Db 120 KYDVYLINPQKAKPRSKVELIAYFEKVDTSIDPNDPFTV-TGRGSPSR 168

RESULT 14
MEC2_RAT STANDARD; PRT; 492 AA.
AC 000566;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)

```

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DE Methyl-CpG-binding protein 2 (MECP-2 protein) (MECP2).
GN Name=MeCP2;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
RC TISSUE=Brain;
RX MEDLINE=92293389; PubMed=1606614;
RA Lewis J.D., Meehan R.R., Henzel W.J., Maurer-Fogy I., Juppensen P.,
RA Klein P., Bird A.;
RT "Purification, sequence, and cellular localization of a novel
RL Cell 69:905-914(1992).
CC -1- FUNCTION: Chromosomal protein that binds to methylated DNA. It can
CC bind specifically to a single methyl-CpG pair. It is not
CC influenced by sequences flanking the methyl-CpGs. Mediates
CC transcriptional repression through interaction with histone
CC deacetylase and the corepressor SIN3A (By similarity).
CC -1- SUBUNIT: Interacts with FMBP3 (By similarity).
CC -1- SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-CpG in the
CC genome.
CC -1- TISSUE SPECIFICITY: Present in all adult somatic tissues tested.
CC -1- SIMILARITY: Contains 2 A+T hook DNA-binding repeats.
CC -1- SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC -----
DR EMBL; M94064; AAA41584.1; -.
DR PIR; A41907; A41907.
DR HSSP; P51608; 10K9.
DR RGD; 3075; MeCP2.
DR InterPro; IPR006337; A+T hook.
DR InterPro; IPR001739; Methyl-CpG_bind.
DR InterPro; IPR001739; Methyl-CpG_bind.
DR Pfam; PF01429; MBD; 1.
DR SMART; SM00384; AT_hook; 2.
DR SMART; SM00391; MBD; 1.
DR PROSITE; PS50982; MBD; 1.
KW Direct protein sequencing; DNA-binding; Nuclear protein; Repeat;
KW Repressor; Transcription regulation.
FT DOMAIN 90 162 MBD.
FT DNA_BIND 185 197 A+T hook 1 (By similarity).
FT DNA_BIND 265 277 A+T hook 2 (By similarity).
FT DOMAIN 366 372 His-rich.
FT DOMAIN 379 403 Pro-rich.
SQ SEQUENCE 492 AA; 53047 MW; A67E705C68BA2D38 CRC64;

Query Match 6.6%; Score 201; DB 1; Length 492;
Best Local Similarity 22.9%; Pred. No. 0.00039;
Matches 108; Conservative 57; Mismatches 156; Indels 150; Gaps 18;

QY 34 LRKE-----DVAMELERVGDEBQMIKRSSECNPLIQEPIASQFG----- 75
   : : : : :
Db 8 LRKEKSDODLQGLKEKPLKFKVKKDKEDKCKHPLPQSAHSAEPBAGAEATSES 67

QY 76 -----ATAGRECKSV-----PCGMEVVKORLFGKTAGRPVYFISP 113
   : : : : :
Db 68 SGSAFAPVPEASASAPKQRRSIIRDRGPMYDPTLPBGWTRKLRKQKSGRSAGKIDVYLINP 127

QY 114 QGLKFRKSSILANYLHNGETSLKPEDPFTVLSKRGKISRKYOCMAALTSHLQNSNN 173
   : : : : :
Db 128 QGKAFRSKVELIAYFEKVDTSIDPNDPFTV-TGRGSPSR----- 167

QY 174 SNNVLRTRSKCKQDVFMPPSSSSSELQSRGLSNFTSHLLKEDGVDVNFPRKV-RKPK 232
   : : : : :
Db 168 -----RQKPPK-----PKSPKAPGTGRGRGRPKSGTGRPKAAASGVQVKKVLEKSP 217

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OY 233 GKVTLKGIPIKTKKGRKSCSGFVQSDS-----KRESVCKNKADESEPVAKSOLDRT 287
DB 218 GK--LLVKKPFGASFGC--KGEGGATTSAGVWVIKRPKPKAKADAPQAIPIKPKR----- 268
OY 288 VQISNAGAGETLSTSENSLVKKKERLSGSGNFCSGPKQTSGLINFKCSAKDSEHNK 347
DB 269 -----GRRKPGSVAAAAAAAKKKAVERSSIR-----SVQETVLPIK-----RK 307
OY 348 YEDTFLSESIGTKYEVVERKEHLTDLL--KRSEMDNNCSPTKDPTEGKIPOEDTTP 405
DB 308 TRET-----VSLFEVAVVPLVSTLTGEGSGKGLTKCKSPGKR-----SKESP 351
OY 406 RTQIERRTKSLYFSKRYKKAALSPRR-----KAFKWTTPRSP 444
DB 352 KGR-----SSSASSPPKKEHHHHHHHAESPKAQMPPLPPPP 388

RESULT 15
MEC2 MOUSE STANDARD; PRT; 484 AA.
ID AC 0923D6;
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Methyl-CpG-binding protein 2 (Mecp2 protein) (Mecp2).
GN Name=Mecp2;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=CS7B1/6;
RC MEDLINE=98449942; PubMed=9774669;
RA Hendrich B., Bird A.;
RT "Identification and characterization of a family of mammalian methyl-
RT CpG binding proteins."
RL Mol. Cell. Biol. 18:6538-6547(1998).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=99299240; PubMed=10369871; DOI=10.1093/nmg/8.7.1253;
RA Coy J.F., Sedlacek Z., Beecher D., Delius H., Fousteka A.;
RT "A complex pattern of evolutionary conservation and alternative
RT polyadenylation within the long 3'-untranslated region of the methyl-
RT CpG-binding protein 2 gene (Mecp2) suggests a regulatory role in gene
RT expression."
RL Hum. Mol. Genet. 8:1253-1262(1999).
RN [3]
RP SEQUENCE FROM N.A.
RA Reichwald K., Thiesen J., Wiehe T., Kloeckle P., Straetling W.H.,
RA Rosenthal A., Platzer M.;
RT "Comparative analysis of the methyl CpG binding protein 2 locus in man
RT and mouse reveals new untranslated sequences."
RT Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RX STRAIN=FVB/N; TISSUE=Mammary gland;
RC MEDLINE=22386257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Udell T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosak S., McMan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulik S.W.,
RA Vallion D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Rahney J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A.C., Shevchenko Y., Bouffard G.G.,
RA Blakeley R.W., Touchman J.W., Green E.D., Dickson M.C.,

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RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield V.S.N., Krzywinski M.I., Skalska U., Smallus D.E.,
RA Schermer A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [5]
RP INTERACTION WITH FMBP3.
RX MEDLINE=97315177; PubMed=9171351; DOI=10.1093/emboj/16.9.2376;
RA Bedford M.T., Chan D.C., Leder P.;
RT "FMBP WW domains and the Abi SH3 domain bind to a specific class of
RT proline-rich ligands."
RL EMBO J. 16:2376-2383(1997).
CC -1- FUNCTION: Chromosomal protein that binds to methylated DNA. It can
CC bind specifically to a single methyl-CpG pair. It is not
CC influenced by sequences flanking the methyl-CpGs. Mediates
CC transcriptional repression through interaction with histone
CC deacetylase and the corepressor SIN3A (By similarity).
CC -1- SUBUNIT: Interacts with FMBP3.
CC -1- SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-CpG in the
CC genome.
CC -1- SIMILARITY: Contains 2 A.T hook DNA-binding repeats.
CC -1- SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; AF072251; AAC68880.1; -
DR EMBL; AJ123922; CAB46495.1; -
DR EMBL; AF121351; -; NOT ANNOTATED_CDS.
DR EMBL; AF158181; AAF33024.1; -
DR EMBL; BC027153; AAH27153.1; -
DR HSPF; P51608; 10K9.
DR MGD; MGI:99918; Mecp2.
DR GO; GO:0005634; C:nucleus; IDA.
DR InterPro; IPR000637; A+T hook.
DR InterPro; IPR001739; Methyl-CpG_bind.
DR Pfam; PF01429; MBD; 1.
DR SMART; SM00384; AT_hook; 2.
DR PROSITE; PS50982; MBD; 1.
DR DNA-binding; Nuclear protein; Repeat; Repressor;
KW Transcription regulation.
FT DOMAIN 90 162 MBD.
FT DNA_BIND 185 197 A.T hook 1 (By similarity).
FT DNA_BIND 265 277 A.T hook 2 (By similarity).
FT DOMAIN 366 372 His-rich.
FT DOMAIN 379 403 Pro-rich.
SQ SEQUENCE 484 AA; 52307 MW; 62FDD28F0118A49F CRC64;

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Query Match 6.5%; Score 198.5; DB 1; Length 484;
Best Local Similarity 24.1%; Pred. No. 0.00054;
Matches 91; Conservative 50; Mismatches 154; Indels 83; Gaps 11;
OY 76 ATAGTBCRSKVS-----PCGMEVYVQRLFGKTAGRPDYVFTSPQGLKFRSKS 122
DB 77 AASAPFQKRSITRDGPMWDTTLPEGWTBKLOKSGRSAGKIVYLINPGKXAFRSKY 136
OY 123 SLANYLHKNGETSLKPEDEFTVLKRGIKSRKYKCSMAALTSILOONSNNSNMILRTS 182
DB 137 ELIAFEBKXGDTSLDNDPDTLV--TGSGSPSRREQ-----KPPKPKSPKAPGTGRGK 189
OY 183 KKKQDVFPSPSSSELQSRGSLNFTSTLLAKEBGVDVNVFRVRRKPKK-----VT 236
DB 190 R-----PKSGTGRPRKAAASBGVQVRRVLEKPGKLVVMPPOASGSGGEGGATT 241
OY 237 ILKGIPIKTKKGRKSCSGFVQSDSKB-----SVCNKADESEPVAKS-----OLDRT 287

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Db	242	SAQVWVIK--RPRKRKAADPPQAI	PKRGRKPGSVAAAAAAEAKKAVKESIRS	VHET	299
Oy	288	VCISDAGACGETLSTSE	-----NSLVKKERSLS	SGSNFCSEOKTSGIINKFC	SAK 340
Db	300	VLPIKKRTRETVSIEVREVV	KPLVSTLGEKSGKGLTKCKSPGRK	SESSPKGRSSSAS	359
Oy	341	DSEHNEKYEDTFLSEEBIGT	KVEVERKEHLHTDILK-----	RGSEMDNNCSP	388
Db	360	SPPKKEHH-----	HHHSESTKAPMPLPSPPP	PEPESSEDPI	ISP 400
Oy	389	TRKDFTEKIFQEDTIPR	406		
Db	401	PEPDLSSSICKERMPR	418		

Search completed: August 22, 2005, 10:09:03
Job time : 182 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:08 ; Search time 42 Seconds
(without alignments)
1328.708 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLESLSLQDRCAPTV.....HKAKYHDMWLNENKLSLS 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: p1r1:*
2: p1r2:*
3: p1r3:*
4: p1r4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	209	6.8	476	2 S57963	methy1 Cpg binding
2	201	6.6	492	2 A41907	methy1-Cpg-binding
3	145	4.7	1301	2 S51323	SAC3 protein - yea
4	145	4.7	3418	1 G02334	breast cancer tumo
5	141.5	4.6	782	2 S27833	thoptry-associated
6	138.5	4.5	500	2 S55785	nucleolar protein
7	137	4.5	560	2 S53382	protein YKR029c ho
8	136.5	4.5	598	2 B40713	cyliclin I - human
9	136.5	4.5	1359	2 T34036	hypothetical prote
10	135	4.4	822	2 T41622	probable ABC trans
11	132.5	4.3	286	2 A72682	probable A/G-spect
12	132	4.3	1819	2 A71928	cag island protein
13	131.5	4.3	997	2 T43523	cut17 protein - fi
14	130	4.3	1927	2 G64585	cag pathogenicity
15	128.5	4.2	946	2 A96748	hypothetical prote
16	128	4.2	853	2 T51505	hypothetical prote
17	126	4.1	561	2 H86442	unknown protein li
18	124.5	4.1	491	2 T50346	hypothetical prote
19	124	4.1	650	2 T33350	hypothetical prote
20	124	4.1	1702	2 T14050	protein kinase (EC
21	123	4.0	697	2 C97120	topoisomerase I (l
22	122.5	4.0	669	2 S55024	nebulin, skeletal
23	122	4.0	651	2 C86333	hypothetical prote
24	122	4.0	3924	2 S37431	ankyrin 2, neuroma
25	120.5	3.9	911	2 S51441	hypothetical prote
26	120.5	3.9	1147	2 JN0599	DNA-binding protei
27	120	3.9	1233	2 S56271	hypothetical prote
28	120	3.9	1131	2 A49393	activator 1 large
29	120	3.9	1440	2 T33813	hypothetical prote

30	120	3.9	1658	2 S55101	hypothetical prote
31	120	3.9	2253	2 T30336	nuclear/mitotic ap
32	119.5	3.9	1740	2 T43773	hypothetical prote
33	119.5	3.9	3122	2 T17202	DNA-directed DNA p
34	119	3.9	646	2 F71620	hypothetical prote
35	119	3.9	797	2 H04919	hypothetical prote
36	119	3.9	991	2 H06168	hypothetical prote
37	119	3.9	1002	2 T30546	major surface glyco
38	118.5	3.9	891	2 B84614	hypothetical prote
39	118.5	3.9	1040	2 E71412	hypothetical prote
40	118	3.9	533	2 B84590	hypothetical prote
41	118	3.9	734	2 B42680	nucleolar-cytoplas
42	118	3.9	1200	2 A46194	neurofilament prot
43	118	3.9	1377	2 T51447	transcription regu
44	118	3.9	3329	2 T42205	breast cancer susc
45	118	3.9	3329	2 T30904	breast cancer tumo

ALIGNMENTS

RESULT 1

S57963 methy1 Cpg binding protein 2 - human (fragment)
C:Species: Homo sapiens (man)
C>Date: 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change 05-Nov-1999
C:Accession: S57963
R:d'Esposito, M.; Quaderi, N.A.; Ciccodicola, A.; Bruni, P.; Esposito, T.; D'Urso, M.; B submitted to the EMBL Data Library, July 1995
A:Description: Physical mapping and expression analysis of an X-linked gene encoding a m
A:Reference number: S57963
A:Accession: S57963
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-476 <DB>
A:Cross-references: EMBL:X89430; NID:g899295; PIDN:CAA61599.1; PID:g899296

Query Match 6.8% Score 209; DB 2; Length 476;
Best Local Similarity 24.0% Pred. No. 6.5e-06;
Matches 104; Conservative 53; Mismatches 149; Indels 128; Gaps 17;

QY	37	EDVAMELEBRVGEDEBQMMIKRSSECNPLLOEPIASAOFG-----ATAGTE	81
DB	13	KDKPLKFKKVKKDKKEKEGHEPVQPSAHSAEPAEAKATSESGSARLCEASAPK	72
QY	82	CRKSV-----PCGERVVKQPLRGKTNGRPDVYFISQGLKFRKSKSLANTL	128
DB	73	QRRSIIIRDGRPMYDPTLPBGWTRKLKORKSGRSAGKYDVYLNPOGKAFRSKVELIAYF	132
QY	129	HONGETSLKPEPFDFTVLSKGIKSRKYDCSMAALTSHLQNSNNNNLFRSKCKKDV	188
DB	133	EKVGDTSIDPNDPFTV-TGRGSPSR-----RKQKPPK--	165
QY	189	FMPSSSSRLQSRGL---SNFTSHLLKEDGVADVFRKVRPKGKVTLLKGIPIK	245
DB	166	---PKSPAPRGGRGRGKRGKSGITTRPKAATSEGVQV--RLLEKSPGK--LLVMQPF-Q	217
QY	246	TKKGRKSCSGFVQSDS---KRESVCKNADSESPVAOKSOLDRTVCISDAGAGETIS	301
DB	218	TPPGKAEGGAGTTSQVWVIRPKRKRAEADPOAIPKR-----GRKGSVVA	267
QY	302	VTSSENSLVKKERLSGSSNFCSEQKTSIINKCSAKDSHNEKYEDTFLESEIGTK	361
DB	266	AAAEKAKKAVESIR-----SVQETVLPFKK-----RTKRT-----VS	303
QY	362	VEVERKEHLHTDIL--KRGSEMNNCSPTRKDFTEKIFQEDTTPRQIERRKTSLYPS	419
DB	304	IEVKEVVKLVYSTLGKSGKGLKTKCKSGRK-----SKESSPKG-----	344
QY	420	SKYNKALSPPRK 433	
DB	345	---SSASAPPKK 355	

A:Accession: S68087
A:Status: nucleic acid sequence not shown; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 374-407 <VAN>
R:Brown, D.; Churcher, C.M.; Barrell, B.G.; Rajandream, M.A.; Wood, V.
submitted to the EMBL Data Library, September 1997
A:Reference number: Z21733
A:Accession: T37634
A:Status: preliminary; translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 1-338, 'S', 340-500 <BRO>
A:Cross-references: EMBL:Z59051, PTDN:CAB11772.1, GSPDB:GN00066, SPDB:SPAC13F5.09
A:Experimental source: strain 972h-; cosmid c13F5
C:Genetics:
A:Gene: gar2; SPDB:SPAC13F5.09
A:Map position: 1
C:Superfamily: ribonucleoprotein repeat homology
F:264-331/Domain: ribonucleoprotein repeat homology <RRM1>
F:367-433/Domain: ribonucleoprotein repeat homology <RRM2>

Query Match	4.5%	Score 138.5	DB 2	Length 500
Best Local Similarity	23.7%	Pred. No. 0.23,		
Match 74	Conservative 50	Mismatches 147	Indels 41	Gaps 11
QY	185	KKDYVMPSSSSSELOESRGLSNFTSTHLLKDEGVDDVDFPKKVPKPKGKVTIILKGIDIK	244	
DB	19	KKGATLEKTSKKKIRKEAKE-----IAQSSKTIDVSPKSKKKEAKRASPE--ISK 68		
QY	245	KTKGCKRKSCSGFVSDSKRESVCKNKADESEPAVQKSOLORTVICISDAGAGETLSTVS	304	
DB	69	KSVKOKKSKK---KESSSESESSSSSSSESSSSSSSSSSSSSSSSSSSSSSSSSSSSSS	122	
QY	305	EENSILVK---KKEBLSGSGNFCSEOKTSGIINKKCSAKDSHNEKYETPLSEBELGTX	361	
DB	123	EEVAVLTKEEKKESSSESSSSSESEEBAAV-KIEEKESSDSSSSSSSSSSSSSSSS	181	
QY	362	VEVVERKHLH-TDLIKRGESEMDNNCSPTKQDFTGEKIFOEETIPIRTOIERRTLTLYSS	420	
DB	182	SESEEEVEVETKEKKGSS	234	
QY	421	KYNKEALSPPRRKAKFKWTPPRSPFNVLVOETLFDHPMKLLIATIFIANRTSGMALPVIWK	480	
DB	235	DEKRRKAPASEERPAKITTPSODSN---ET-----CTVAVGRISNVWDQWIGQ	281	
QY	481	FLEKYPASAVAR 492		
DB	282	EFEETGTVGAR 293		

RESULT 7
 S53382
 protein YKR029C homolog YUL105w - yeast (*Saccharomyces cerevisiae*)
 N/Alternate names: hypothetical protein J0819
 C/Species: *Saccharomyces cerevisiae*
 C/Date: 05-May-1995 #sequence revision 01-Sep-1995 #text_change 09-Jul-2004
 C/Accession: S53382; S56883; S57363
 R/Rasmussen, S.W.
 submitted to the EMBL Data Library, February 1995
 A/Description: A 37.5 kb region of yeast chromosome X includes the SWE1, MEF2, GSH1 and
 A/Reference number: S53376
 A/Accession: S53382
 A/Molecule type: DNA
 A/Residues: 1-560 <RAS>
 A/Cross-references: UNIPROT:P42948; EMBL:X85021; NID:g728698; PID:g728705
 R/Rasmussen, S.W.
 submitted to the Protein Sequence Database, September 1995
 A/Reference number: S56876
 A/Accession: S56883
 A/Molecule type: DNA
 A/Residues: 1-560 <RAW>
 A/Cross-references: EMBL:Z49380; NID:g1008285; PID:g1008286; MIPS:YUL105w
 R/Rasmussen, S.W.
 yeast_11, 873-883, 1995

A1Title: A 37.7-kb region of yeast chromosome X includes the SWR1, MEF2, GSH1 and CSD3 genes
A1Reference: number: S57357; MUID:96090136; PMID:7483851
A1Accession: S57363
A1Status: nucleic acid sequence not shown; translation not shown
A1Molecule type: DNA
A1Residues: 1-360 <RAN>
A1Cross-references: EMBL:X85021, NID:g728668; PIDN:CA59389.1; PID:g728705
A1Note: The nucleotide sequence was submitted to the EMBL data library, February 1995
C1Genetics:
A1Cross-references: SGD:S0003641
A1Map position: 10L

	Query Match	4.5%	Score 137;	DB 2;	Length 560;
	Best Local Similarity	18.7%	Pred. No. 0.33;		
	Matches	90;	Conservative	80;	Mismatches 161; Indels 150; Gaps 24
QY	118 FRKSKSLANYLHKNGETSLKEPEDDPFVLSKRGISRKRCDCSMALTSHTL--QNQSNNSN	175			
Dd	82 FQNRGRIINHHSSSGSSKT-----ASTNKKRGIAAAVALATAATIPPLKKNQDDNSK	134			
QY	176 WNLRTSRCKCKDVFEMPPSSSELQESRGLSNFTSTHLLIKEDGVDDYNFRKKRPKGY	235			
Dd	135 VSV-THNESSEKENKITTPSMAE-----DNKPKNGCICGSSDSXDEL-----	174			
QY	236 TILMGIPITKTKRKGCRKSCGFVSDS-KRSVCNKADAESEPVAOKSQLD-----	285			
Dd	175 ----PIQCNKKCTWOHQLCYAFPKSDPIKRDVCAKCDSDTK--YQNVQVKMIFPRKMG	228			
QY	286 -----RTVCISPAAGCE-----TLVSISEENSLVK-KERSLS	318			
Dd	229 DERLFQFSIYTTSANTNHQGSVNNIIEQPKRQHLYTAPTTENSINSIRKLARQEKLV	288			
QY	319 SGNFCSFQGTSGIINFCSAKD-----SEHNKEDTFLESEEIGTKVVEVERKH	370			
Dd	289 VSSHFLKP-----LNLEVSSNDIEFKAITSEYDKCVKVFIDN-----H	329			
QY	371 LHTLIL-----KRGEMDNCCSPTRDPTFGBKIFQEDLIPTRO-IRRKTSLEYSSKY	422			
Dd	330 YDDMVAVVCSNMESSASADIIVKKSNEBDF--GFPAADS CVKGELIOBYLAKIDPOKNY	386			
QY	423 NKEALSPPRRKAFKMTKTPPSPPFNVLQET-----LPHDWPKLLATIIFLNRTSGKMAIPVL	478			
Dd	387 Q-----IDPNNDYLMLGWTTRKVLFLHPMPLYDS--REHG--L	422			
QY	479 WKLETG--YPSABAVARTADMWDVSELLKPLGLYDIRAITYKF-----SDEYLLIKOMKYP	531			
Dd	423 TRYLRSCPEPVELVTY--RPLDS--KPRGDNCRAVFVRLAIRIDIRKGEIISVEMQWD	477			
QY	532 I 532				
Dd	478 L 478				

```

RESULT 8
B40713
CylC1c1n I - human (fragment)
C|Species: Homo sapiens (man)
C|Date: 12-May-1994 #sequence_revision 12-May-1994 #text_change 09-Jul-2004
C|Accession: B40713; S35920
R|Hess, H.; Heid, H.; Franke, W.W.
J. Cell Biol. 122, 1043-1052, 1993
A|Title: Molecular characterization of mammalian cylc1n, a basic protein of the sperm he
A|Reference number: A40713; MUID:93359502; PMID:9354692
A|Accession: B40713
A|Status: Preliminary
A|Molecule type: mRNA
A|Residues: 1-598 <HES>
A|Cross-references: UNIPROT:P35663; GB:Z22780; NID:g396104; PIRN:CAA80457.1; PID:g396105
C|Keywords: cytoskeleton

Query Match          4.5%; Score 136.5; DB 2; Length 598;
Best Local Similarity 23.3%; Pred. No. 0.39;
Matches 111; Conservative 72; Mismatches 205; Indels 89; Gaps 21;

```


QY 2 GTTGLESISLGDGGAAPVTVTSSERLVPDPNDLRKEDVAMELERVGEDEBQMIRSSSEC 61
 DB 70 GGTPLKPKDSKKKGGSVATPBEKQIVBEKTKRQONAD- KTLPLKSHENEQSKKSSSET 128
 QY 62 NPLOEPASAOFGATAGTECKSVPCGMEVVRVORLFGKTAGRDVAFI-----SPOGLK 117
 DB 129 NP-----ESQNSKTVSKNCSQ-----KOKOSKSKKKTNTTEFLTKNNPKODL 171
 QY 118 FSKSKSLANYLHKNETSLKPEDPFTVLSKRGISRYKDCSMAALTSHLQON-OSNSNN 175
 DB 172 KRSKTSNDPISICENSJ-NDPFLMVGQ-----SDDESINPDMALRYSQNSGN 222
 QY 176 WNLRTSRCKCKDV-FMPSSSSELQESR-GLSNTSTHLLKEDBGVDV----- 223
 DB 223 YELKTKTKTKODTKKNAKSSDAESEDSDKAKKSKVKNVKKODKKKDVKKDTESTDA 282
 QY 224 -----NPKVAKPKGKVTILKGIPIKTKK-----GCRKSCSGFVQS 260
 DB 283 ESGDKDERKDYK-KDKKKLKDDKKDKDYKYPSTDTESGDAKARDNSRLKKAASKND 341
 QY 261 DSKRES--VCNKADAESEFPAQSOQLDRTVCISDAGCETLSVTEENSLVKKKERSLS 318
 DB 342 DKKKAKKITTSTDBSELESSEKSOQDEKKDD-----SKTDNKKK-VKNDEESTD 392
 QY 319 SGSNFCSEQKTSGLINKFCSAKDSEHNEYEDT-FLSEELGTQVVEVERKEHLHTDIL 376
 DB 393 ANSEPKGDSK-KGKDEKKKKDSKKDKKDAKNAKSTEMSDLELKKDKKHSKE--- 448
 QY 377 KRGSEMDNNSPFRKDFTEKLFQEBTTRTOIERKTKSLYSSKTKNALSPPRK 433
 DB 449 KKGSKKDKIK-KDARKD-TESTDAFDESSKTGF-KTSTIKGSDTESESLYKPKAK 502

RESULT 9

T34036
 Hypothetical protein B0041.7 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 09-Jul-2004
 C:Accession: T34036
 R:Fullon, R.; Wohlmann, P.
 Submitted to the EMBL Data Library, April 1997
 A:Description: The sequence of C. elegans cosmid B0041.
 A:Reference number: Z21466
 A:Accession: T34036
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-1359 <FUT>
 A:Cross-references: UNIPROT:Q9U7B0; EMBL:AF000196; P1DN:AAC24256.1; GSPDB:GN00019; CESP:
 A:Experimental source: strain Bristol N2; clone B0041
 C:Genetics:
 A:Gene: CSP; B0041.7
 A:Map position: 1
 A:Introns: 12/2; 59/2; 248/2; 582/1; 675/1; 733/3; 966/2; 1044/2; 1310/3

Query Match 4.5%; Score 136.5; DB 2; Length 1359;

Best Local Similarity 19.2%; Pred. No. 1.2;

Matches 113; Conservative 91; Mismatches 253; Indels 131; Gaps 19;

QY 15 GAAPVTSSERLVPDPNDLRKEDVAMELERVGEDEBQMIRSSSECNPLLOEPASAO 74
 DB 4 GVESESDSDGHVIED-EDL-EMARQIENRERKKAQKLEKRE----- 44
 QY 75 GATAGTECKSVPCGMEVVRVORLFGKTAGRDVAFI-----SDDESINPDMALRYSQNSGN 134
 DB 45 -----RSGKPPPKRPAKRRKASSSEDDDEESPR--KSSKSKRRKASSSESD 94
 QY 135 SLKPEDPFTVLSKRGISRYKDCSMAALTSHLQONNSNNWNLRTSRCKCKDVFMPPSS 194
 DB 95 SDEBEDPKSKSKKKVQKKKKSKKRTSSSEDESDEREGKSKKKSKTKKQOSSE 154
 QY 195 SSELQESRGLSNTSTHLLKEDBGVDVNFRTYRK-PKGVITLTK-----GI 241

DB 155 SSEESEB-----RKVKSKKONKESKVKRAETSESDDEK 191
 QY 242 PIKTKKGCRCRSGGFVQSDSKRESVCNKADAESEFPAQSOQLDRTVCISDAGCETLS 301
 DB 192 PSKSKKGLKRAVSESESEDEKEVYKSKKKKVVKKS-----SEEDNAPEKK 244
 QY 302 VTSEENSLVKKKERSLSGNSFCSEOKTSG-----INKFCSAKDSHNEYEDTF 352
 DB 245 TEKKRSTSTSESESESKDDEEBEKSSPKPKKKPLAVAKULSDSESD- 298
 QY 353 LESEELGTQVVEVERKEHLHTDILKRGSEMDNNSPFRKDFTEKLFQ-EDTTRTOIER 410
 DB 299 -----VEVLPOK-----KRGAVTILISDEBEKDKKSSSEASDVEEKVSKKAK 342
 QY 411 RRTSLVSSKTKNEALSPPRKAFKWTTPRSPFNIVQETLPHDPWTLATIFLNRTS 470
 DB 343 KQES-BSGSDSGSITVNRKSKKKKKKKGIIM-----DSKLOKETIDAEAE 395
 QY 471 GKMAIPVLMKFLK-KYPSAEVARTADMRDVELLPLGLYDLRAKTIKVFDEVLTKQWK 529
 DB 396 KERR-----KLEKKQKFPNGIVLEGEDLTMLTGTSQKRLKSVLDPPSSTVDESK 450
 QY 530 YPIELHGIQKYGNDSYRIFCVNEMKQVPEHDKLNKYHDMWENHEKL 577
 DB 451 KPVEVH-----NSLVRL-----KPH-QAHGIQFMWDCAFEESIDRL 485

RESULT 10

T41622

probable ABC transporter - fission yeast (Schizosaccharomyces pombe)

C:Species: Schizosaccharomyces pombe

C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 09-Jul-2004

C:Accession: T41622

R:Aert, R.; Voickaert, G.; McDougall, R.C.; Rajandream, M.A.; Barrell, B.G.

Submitted to the EMBL Data Library, October 1999

A:Reference number: Z21735

A:Accession: T41622

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-822 <AER>

A:Cross-references: UNIPROT:Q9USH9; EMBL:AL122011; P1DN:CAB58409.1; GSPDB:GN00068; SPDB:

A:Experimental source: strain 972h-; cosmid 6825

C:Genetics:

A:Gene: SPDB:SPCC825.01

A:Map position: 3

Query Match 4.4%; Score 135; DB 2; Length 822;

Best Local Similarity 22.4%; Pred. No. 0.75;

Matches 80; Conservative 53; Mismatches 134; Indels 90; Gaps 14;

QY 149 RGIKS--RYKDCSMAALTSHLQONNSNNWNLRTSRCKCKDVFMPPSSSELQESRGLSN 206
 DB 3 RGRSTQRADULESLQESIESPQP-----VTRSKAK-----NKKKLNKASAFNS 51
 QY 207 FSTHLLK-EBEGVDVNFRTYRKPKGKVTILKGIPIKTKKGCRCRSGGFVQSDSKRE 265
 DB 52 DNDSDYDLKPEDDEVD-----EVVVKKKPKSKKKAANAEAPADEQSV 100
 QY 266 SVCNKADAE-----SEFPAQSOQLDRTVCISDAGCETLSVTEENSLVKKKERS 316
 DB 101 E--EEDSKPVRKAKSKSKASPKNAFALDMDLSDLEBESSSSKSKKKKKKSKS 158
 QY 317 LSGSNFCSEOKTSGIINKFCSAKDSEHNEYEDTFLESEELGTQVVEVERKEH---LH 372
 DB 159 KDDG-----SEALDDGI-----ESSEKKKKKKKSKENDDAFKDKRTRKKEEKARAKLA 208
 QY 373 TDILKRGSEMDNNSPFRKDFTEKLFQEBTTRTOIERKTKSL 416
 DB 209 SMLESENKNDANAAALNTTDAFKDGLPGRLIFAVASQCAVAPDSNADGI----- 261
 QY 417 YFSKTKNEALSPPRKAFKWTTPRSPFNIVQETLPHDPW-KLITATIFLNRTSGK 472
 DB 262 ----TGTGMLSPNSR-----DLQVKLSVAMGWLILKQSELMNLINCR 302

174 SWNNITRSKCKKD-----VFMPSSSSSEIQES-----RGLSNFTSTHLLKEDEGVD 222
 587 SEOSIQLESSEBNDKPLDLPILAIRKKNLVGVLEKKGSTSTSK---TKFTSTLYD 643
 223 VNFVRKVRKGVTLIKGPIKTKKCKGKSGGFVQ-----SDSKRESV 267
 644 F-----IEKPTKIS--EVLPEKRAKICDESQTVRVSIDRGVTKTRDVSPPISDEKSENV 697
 268 CNKADAES-----BPVAKSQDLRTVCISDA--GACGETLSTVSEENSLVVK 312
 698 -NHEEANGSHVTWNVHSSLDPOPIVQPNBELSGSYLKLDPRNVGNSEKVTFOEDDINS 756
 313 KERSJSSGNSFCSEOKTSGIINKFCGAKSENEKEDPFLSEELGTAVVEKREH 372
 757 KIQSKNQVNAVNTSETSDKQEKEMHLENIKEKLEKTEVDKSLSDAPDEQIKNS 816
 373 TDLKRGSEMDNNSCPTKDFGKIFQEDTIPTQIER-----KTSLYSSKY-NKEAL 427
 817 RTSVQNGTSSVSCKMTPKE-----TKVDKIDNVSKDQVETSPGSCETISAFATYAEKVT 872
 428 S---PPRRKAFYK--WTPRSPFNLVQETLPHDPWKLITATILNRTSGMAIPVLMKFL 482
 873 SINDLSVRKPLDESYVDHISFPDPLCGSSFLAP-----QFPVSKKHALPLV----- 919
 483 EKYPSAEVARTDWR--DVSELK-----PLGLYDLRACTIVKFSDEYLTQW-KPIPL 534
 920 -----EANAPWEPIDFSSLESFVNPVBNKLSKEL-----DMTVEQMIK----- 963
 535 HGIGKYGNSYRIFCNEMKQVHPEDHKKNKHDMLMENHEK 576
 964 ---MYAK-----CAKEFEACEE--KI-----EWLESGKR 989

RESULT 14

66585
 cag pathogenicity island protein cag7 - Helicobacter pylori (strain 26695)
 C1Species: Helicobacter pylori
 C1Date: 09-Aug-1997 #sequence_revision 09-Aug-1997 #text_change 09-Jul-2004
 C1Accession: G64585
 R1From: J.F. White, O. Kerlavage, A.R. Clayton, R.A. Sutton, G.G. Fleischmann, R.D. Peterson, S. Loftus, B. Richardson, D. Dodson, R. Khakh, H.G. Glodek, A. McKenney, J.D. Kelley, J.M. Cotton, M.D. Weidman, J.M. Fujii, C. Bowman, C. Matthey, L. Nature 388, 539-547, 1997
 A1Authors: Wallin, E.; Hayes, W.S.; Borodovsky, M.; Karpk, P.D.; Smith, H.O.; Fraser, C. A1Title: The complete genome sequence of the gastric pathogen Helicobacter pylori.
 A1Reference number: A64520; MUID:97394467; PMID:9252185
 A1Accession: G64585
 A1Status: preliminary; nucleic acid sequence not shown; translation not shown
 A1Molecule type: DNA
 A1Residues: 1-1927 <TOM>
 A1Cross-references: UNIPROT:Q25262; GB:AE000567; GB:AE000511; NID:92313641; PIDN:AA0759

Query Match 4.3%; Score 130; DB 2; Length 1927;
 Best Local Similarity 20.5%; Pred. No. 5;
 Matches 89; Conservative 61; Mismatches 147; Indels 138; Gaps 21;
 17 APTTSSERLVPPNDLRKEDVAME-----LRRVGEDEBQMMIKRSSGPNLL---OB 67
 1137 AKTEAKKKGKCVKLPDLOKVKYAKSVAYLDCVBARNE--KKKKECKLLTPEAKK 1193
 68 PIASAOFGATAGTECKSVPCGMEVYVKKRLFEKTAGR-----PDVY----- 109
 1194 LLEAKESLKAAYDCLSQANBEERACEKLLTPEARKLLEQEVKKSIKAYLDCVBARN 1253
 110 -----FISPOGLKFRSKSLAYLHNG-----ETSLK--PRDPFTVLSKRGIS 153
 1254 EKEKKECKLLTPEARKFLAKOVL-NCLEPAQNEBERKACKLKNLPDLOENILAKSLSVA 1312
 154 RYDCGMAALTSHLQONSNNSNNLRTSRCKCKDVMPSPSSSELOESRGLSNFTSTHLL 213
 1313 -YKDC-----LSGARMBEE-----RRGCEK--LLTPEARKLLEQ----- 1343
 214 LKEDEGVADVNFRRKVRK--KGKVTILKGIPIKTKKCGKRSKSGGFVQSDSKRESVCKND 272

1344 -----EYKSVKAYLDCVSAARBEKKECKELT----- 1373
 273 AESEVPAK-----SQDLRTV--CI-----SDAGACETLSTVSEENSLVKKERSLSS 319
 1374 -----PEARKFLAKEIQCDKAIKDCLKNADBNDAATIKCDLGSDEKEL----- 1419
 320 GSNFSEOKTSGIINKFCGSAKDSSENEKEDPFLSEELGTAVVEKREH 378
 1420 --KYLQERKAVNDCAMATDEKRRKQNVSLDIOENKTKONKONLSTERLHQ 1477
 379 GSE-MDNNSCPTK 392
 1478 ASECLINDDPTDQE 1492

RESULT 15

A96748
 hypothetical protein T10D10.13 [imported] - Arabidopsis thaliana
 C1Species: Arabidopsis thaliana (mouse-ear cress)
 C1Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 09-Jul-2004
 C1Accession: A96748
 R1Theologis, A.; Becker, J.R.; Palm, C.J.; Federpiet, N.A.; Kaul, S.; White, O.; Alonso, Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasey, T.H.; Dewar, K.; Jensen, N.F.; Hughes, B.; Huizar, L.
 Nature 408, 816-820, 2000
 A1Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Kim, C. C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Malt, R.; Matzali, R.; Rooney, T.; Rowley, D.; Sakano, H.
 R1Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallon, ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
 A1Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
 A1Reference number: A6141; MUID:21016719; PMID:1110712
 A1Status: preliminary
 A1Molecule type: DNA
 A1Residues: 1-946 <STO>
 A1Cross-references: UNIPROT:O9C9D8; GB:AE005173; NID:96730761; PIDN:AA07150.1; GSPDB:GN C1GeneID:8
 A1Gene: T10D10.13
 A1Map position: 1

Query Match 4.2%; Score 128.5; DB 2; Length 946;
 Best Local Similarity 21.0%; Pred. No. 2.4; Indels 121; Gaps 16;
 Matches 94; Conservative 56; Mismatches 177;
 32 NDLRKEDVAMELERVGEDEBQMMIKRSSGPNLLQEPASQFGATAGTECKSVPCG-- 89
 370 NELELTFAPHQARVCGDSSSRRGKSSSENVAAHPEPHS-----IAATEKRLSLGGGSA 424
 90 -WERYVKORLFGKTAGRPDVYFISFPOGLKFRSKSLAYLHNGETSLKPEDPFTVLSK 148
 425 DESKMTPLVGDKGDALRRNLSDLSLDSKGFYKRYKMKRDAKRE----- 475
 149 RKIKRKYDCGMAALTSHLQONSNNSNNLRTSRCKCKDVMPSPSSSELOESRGLSNFT 208
 476 WSLKGEKRETKKSKQBALEQSRTEMKAKLASSEBRKDL-----STRQAEKFRSN 530
 209 STHLLK-----EDEGVADVNF----- 226
 531 SSSMKKTKQSEBDEDSIEKPPAKKKAASGQSVGSISRSSQARKLPNRMSSSITPR 590
 227 ---KYRKPKGVTLIKGPIKTKKCGKRSKSGGFVQSDSKRESVCKNDASEPVAQSQ 283
 591 SAASVPKPGKVS-----NTSSGRRRS-----DKSLAQSVNFSLEIKENTKPSLS 636
 284 LDRYTCISDAGCGEGLSTVSEENSLVKKKE--RSLSSG-----SNFSEOKTSGIIN 334
 637 AVKTTMRQSVKSGGRTKN--KEDTLQRPRLKRSSSSNIDFTSLTSLCSDM--WVS 691
 335 KFCGSAKDS--ENEKEDPFLSESEI--GTKYVVERKEKHLATDILKRGSEM----- 382
 692 LAVDSDISLTLNNEYEDBEPAPEVLENAVAEEVEVELTFLVPEDGPMLSAEVAKVD 751

QY 383 ---DNNCSPTKRDFTGEXI|FOEDTIPRT 407
: ||| |:
Db 752 HSGENCs-----FLPATVPTT 768

Search completed: August 22, 2005, 10:06:00
Job time : 46 secs

GenCore version 5.1.6
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OM protein - protein search, using SW model

Run on: August 22, 2005, 10:02:08 ; Search time 170 Seconds
(without alignment)
1319.535 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055

Sequence: 1 MGTGLESLSIGDRGAPTV.....HKLNKTHMLWENHEKLSLS 580

Scoring table: BLOSUM62

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database :

1: Geneseq_16Dec04:*
2: geneseqp19808:*
3: geneseqp19908:*
4: geneseqp20008:*
5: geneseqp20018:*
6: geneseqp20038:*
7: geneseqp20038:*
8: geneseqp20048:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3055	100.0	580	2	AAW74473
2	3055	100.0	580	7	ADDB9906
3	3055	100.0	580	8	ADK13954
4	2958.5	96.8	565	3	AAV44504
5	2194.5	71.8	439	2	AAV76548
6	1639	53.6	307	7	ADDB9915
7	1230	40.3	257	2	AAW88701
8	1230	40.3	257	6	ABBS0468
9	1230	40.3	257	6	ABO44725
10	1230	40.3	257	7	ABO26205
11	1106	36.2	202	7	ADDB9916
12	880.5	28.8	416	7	ADDB9908
13	816	26.7	147	7	ADDB9917
14	357	11.7	68	5	AAE22568
15	254	8.3	50	4	ABBS1147
16	254	8.3	50	6	ABO45404
17	254	8.3	50	7	ABO26884
18	220.5	7.2	467	8	ADK13968
19	220.5	7.2	467	8	ADK13967
20	213.5	7.0	466	8	ADK13972
21	209	6.8	476	8	ADJ68460
22	209	6.8	476	8	ADK13973
23	208.5	6.8	477	8	ADK13961
24	208.5	6.8	466	8	ADK13966
25	208.5	6.8	466	8	ADK13971

26	208.5	6.8	486	8	ADK13957	Adk13957 Human met
27	208.5	6.8	486	8	ADK13960	Adk13960 Human met
28	208.5	6.8	486	8	ADK13953	Adk13953 Human met
29	208.5	6.8	486	8	ADK13963	Adk13963 Human met
30	208.5	6.8	486	8	ADK13970	Adk13970 Human met
31	208.5	6.8	486	8	ADK13958	Adk13958 Human met
32	208.5	6.8	486	8	ABM82465	Abm82465 Tumour-as
33	208.5	6.8	560	6	ABG72561	Abg72561 TAR dwt-M
34	208.5	6.8	561	6	ABG72557	Abg72557 Mecp2-Tat
35	202.5	6.6	345	8	ADK14010	Adk14010 Chicken m
36	201	6.6	492	8	ADK13964	Adk13964 Rat methy
37	198.5	6.5	484	8	ADK13959	Adk13959 Mouse met
38	198.5	6.5	484	8	ADK13956	Adk13956 Mouse met
39	198.5	6.5	484	8	ADK13969	Adk13969 Mouse met
40	198.5	6.5	484	8	ADK13965	Adk13965 Mouse met
41	194	6.4	219	3	AAQ02051	Aaq02051 Human sec
42	186	6.1	68	5	AAE22566	Aae22566 Human Mec
43	186	6.1	68	5	AAE22566	Aae22566 Human Mec
44	159.5	5.2	744	8	ADP49805	Adp49805 Human 193
45	148.5	4.9	326	3	AAQ16859	Aaq16859 Arabidops

ALIGNMENTS

RESULT 1
AAW74473 standard; protein; 580 AA.
ID AAW74473;
AC AAW74473;
XX 19-MAY-1999 (first entry)
DT 19-MAY-1999 (first entry)
XX Human MED1 endonuclease protein sequence.
DE Endonuclease; MED1; human; methyl-CpG binding endonuclease-1;
XX DNA fidelity; DNA manipulation; cancer; fragile X syndrome; therapy;
XX myotonic dystrophy; Huntington's disease; spinocerebellar ataxia;
XX Kennedy's disease; triplet repeat expansion disorder.
XX Homo sapiens.
XX OS
XX WO9904626-A1.
XX PD
XX 04-FEB-1999.
XX PF
XX 28-JUL-1998; 98WO-US015828.
XX PR
XX 28-JUL-1997; 97US-0053936P.
XX (FOXC-) FOX CHASE CANCER CENT.
XX Bellacosa A;
XX MPI: 1999-142462/12.
XX N-PSDB; AAX22002.
XX New nucleic acid encoding human endonuclease MED1 involved in DNA
XX mismatch repair - used for diagnosing susceptibility to cancer and
XX fragile X syndrome, and therapeutically.
XX Claim 8; Fig 3; 109pp; English.
XX PS
XX This sequence is the human MED1 endonuclease of the invention. MED1 (for
XX methyl-CpG binding endonuclease-1) is used to screen for specific
XX modulators (potential therapeutic agents particularly mimetics of MED1)
XX and to study interactions involved in maintaining DNA fidelity, for DNA
XX manipulation and to raise antibodies. Susceptibility or predisposition to
XX cancer (particularly colorectal or endometrial, especially hereditary non
XX -polyposis colorectal cancer), or its prognosis, where caused by
XX alterations in the MED1-encoding gene, are identified by sequence
XX comparison, amplification, detecting altered polypeptide, and restriction
XX fragment mapping, hybridisation (particularly to probes specific for a

CC mutant allele). These same methods can also be used to diagnose fragile X
CC syndrome and other diseases (e.g. myotonic dystrophy, Huntington's
CC disease, spinocerebellar ataxia and Kennedy's disease), associated with
CC triplet repeat expansion. The DNA, or its fragments, are used as probes
CC and primers in the above diagnostic methods, also to isolate homologous
CC sequences, as sources of antisense sequences and for gene transfer,
CC particularly to restore drug sensitivity to drug-resistant cancer cells
XX

Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 2; Length 580;
Best Local Similarity 100.0%; Pred. No. 7,7e-287;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```
QY 1 MGTGLESLSIGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
DB 1 MGTGLESLSIGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
QY 61 CNPLLOEPISAQAGTAGTECRKSVPCGWERVVKQRLFKTAGRFDYVFIISPOGLKFRS 120
DB 61 CNPLLOEPISAQAGTAGTECRKSVPCGWERVVKQRLFKTAGRFDYVFIISPOGLKFRS 120
QY 121 KSSLANYLHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLR 180
DB 121 KSSLANYLHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLR 180
QY 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTYILKG 240
DB 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTYILKG 240
QY 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
DB 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
QY 301 SVTSEENSLVKKKERSLSGSGNFCSGEOKTSGIINFKCSAKDSEHNEKYEJDTPLSESEIGT 360
DB 301 SVTSEENSLVKKKERSLSGSGNFCSGEOKTSGIINFKCSAKDSEHNEKYEJDTPLSESEIGT 360
QY 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKIFQEDTTPRTQIERRKTSLYFSS 420
DB 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKIFQEDTTPRTQIERRKTSLYFSS 420
QY 421 KYNKEALSPRRKAFKWTTPRSPFNLYOETLFHDPMLLLATIFLNTSGKMAIPVLMK 480
DB 421 KYNKEALSPRRKAFKWTTPRSPFNLYOETLFHDPMLLLATIFLNTSGKMAIPVLMK 480
QY 481 FLEKTPSAEVARATADWRDVSELKPLGLYDLRAKTIKVFSDSEYLTQWKYPIELHIGIKY 540
DB 481 FLEKTPSAEVARATADWRDVSELKPLGLYDLRAKTIKVFSDSEYLTQWKYPIELHIGIKY 540
QY 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDLWENHEKLSLS 580
DB 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDLWENHEKLSLS 580

RESULT 2
ADD89906
ID ADD89906 standard; protein; 580 AA.
XX
AC ADD89906;
XX
DT 29-JAN-2004 (first entry)
XX
DE Human 5-methylcytosine DNA glycosylase.
XX
KW Human; 5-methylcytosine DNA glycosylase; enzyme; Cpg.
OS Homo sapiens.
XX
PN WO2003078593-A2.
XX
PD 25-SEP-2003.
```

PF 14-MAR-2003; 2003WO-US007933.
XX
XX 15-MAR-2002; 2002US-0364689P.
XX
XX (EPIG-) EPIGENOMICS AG.
PA
PI Lofton-Day CE, Day JK;
XX
XX WPI: 2003-779127/73.
DR
DR N-PSDB; ADD89905.
XX
XX
PT Labeling methylated or methylatable Cpg sequences, useful e.g. for
PT diagnostic detection of altered methylation, comprises replacing
PT methylated cytosine by labeled cytosine.

Claim 11; Page 53-55; 73pp; English.

CC The present sequence is the protein sequence of human 5-methylcytosine
CC DNA glycosylase (5-MCDG). The enzyme acts by cleaving glycosyllic bonds at
CC methylated Cpg sites of DNA, removing 5-methylcytosine from the DNA
CC backbone as a free base. Human 5-MCDG can be used in a claimed method for
CC labelling Cpg sequences corresponding to methylated Cpg sequences in an
CC isolated DNA sample. The method comprises: digesting the genomic DNA with
CC a restriction endonuclease to produce genomic DNA fragments; treating the
CC genomic DNA fragments with 5-MCDG such that one or more 5-methylcytosine
CC bases are removed to produce abasic genomic DNA fragments; and treating
CC these abasic genomic DNA fragments with base excision repair enzymes in
CC the presence of labelled dCTP such that 5-methylcytosine removed from the
CC genomic DNA fragments by 5-MCDG is replaced by labelled cytosine in the
CC one or more corresponding positions of the abasic genomic DNA fragments
CC to produce labelled genomic DNA fragments, so that specific labelling of
CC Cpg sequences corresponding to methylated Cpg sequences is achieved. The
CC 5-MCDG is also used in a claimed method for comparing Cpg methylation
CC status, extent or pattern between or among reference and test genomic DNA
CC samples, and in a claimed method for labelling potentially-methylatable
CC Cpg sequences in Cpg-containing genomic DNA fragments. The methods are
CC used to identify methylated and/or potentially methylatable Cpg
CC dinucleotides in genomic DNA, including comparison of methylation pattern
CC between healthy and diseased samples, for diagnosis.

Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 7; Length 580;
Best Local Similarity 100.0%; Pred. No. 7,7e-287;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```
QY 1 MGTGLESLSIGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
DB 1 MGTGLESLSIGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEQMMIKRSSE 60
QY 61 CNPLLOEPISAQAGTAGTECRKSVPCGWERVVKQRLFKTAGRFDYVFIISPOGLKFRS 120
DB 61 CNPLLOEPISAQAGTAGTECRKSVPCGWERVVKQRLFKTAGRFDYVFIISPOGLKFRS 120
QY 121 KSSLANYLHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLR 180
DB 121 KSSLANYLHKNGETSLKPEDDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSNMNLR 180
QY 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTYILKG 240
DB 181 RSKCKDVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKRGKVTYILKG 240
QY 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
DB 241 IPIKTKKGRKSCSGFVQSDSKRESVCKADASEPVAOKSOLDRTVCISDAGACGRTL 300
QY 301 SVTSEENSLVKKKERSLSGSGNFCSGEOKTSGIINFKCSAKDSEHNEKYEJDTPLSESEIGT 360
DB 301 SVTSEENSLVKKKERSLSGSGNFCSGEOKTSGIINFKCSAKDSEHNEKYEJDTPLSESEIGT 360
QY 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKIFQEDTTPRTQIERRKTSLYFSS 420
DB 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFGEEKIFQEDTTPRTQIERRKTSLYFSS 420
```

QY 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLIIATITFLNRTSGMAIPVLMK 480
 DB 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLIIATITFLNRTSGMAIPVLMK 480
 QY 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 DB 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 QY 541 GNDSTRICVNMKQVHPEDHKLANKYHDMWMENHEKLSLS 580
 DB 541 GNDSTRICVNMKQVHPEDHKLANKYHDMWMENHEKLSLS 580

RESULT 3

ADK13954
 ID ADK13954 standard; protein; 580 AA.

ADK13954;
 AC ADK13954;

DT 03-JUN-2004 (first entry)
 XX

DE Human methyl-CpG-binding protein #9.
 XX

XX Rett syndrome; methyl-CpG-binding protein 2; MECP2;
 KM neurodevelopmental disease; autism; non-syndromic mental retardation;
 KM idiopathic neonatal encephalopathy; idiopathic infantile spasm;
 KM idiopathic cerebral palsy; Angelman syndrome; schizophrenia; human.
 XX

OS Homo sapiens.
 XX

XX US6709817-B1.
 XX

XX 23-MAR-2004.
 XX

XX 07-SEP-2000; 2000US-00657013.
 XX

XX 07-SEP-1999; 99US-0152778P.
 XX

XX (BAYU) BAYLOR COLLEGE MEDICINE.
 XX

XX Zoghbi HY, Van Den Veyver IB, Amir R, Francke U;
 XX

XX MPI; 2004-256068/24.
 XX

PT Screening human for Rett syndrome comprises detecting mutation in nucleic
 acid sequence encoding methyl-CpG-binding protein 2 (MECP2).
 XX

PS Disclosure; SEQ ID NO 56; 125pp; English.
 XX

CC The invention relates to a method of screening a human for Rett syndrome
 comprising detecting a mutation in a nucleic acid sequence encoding
 methyl-CpG-binding protein 2 (MECP2). The method is useful for screening
 a human for Rett syndrome. The method is useful for screening
 CC neurodevelopmental diseases such as Rett syndrome, autism, non-syndromic
 CC mental retardation, idiopathic neonatal encephalopathy, idiopathic
 CC infantile spasms, idiopathic cerebral palsy, Angelman syndrome and
 CC schizophrenia. The present sequence represents the amino acid sequence of
 CC a methyl-CpG-binding protein.
 XX

XX Sequence 580 AA;
 SQ

Query Match 100.0%; Score 3055; DB 8; Length 580;
 Best Local Similarity 100.0%; Pred. No. 7,7e-287;
 Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MGTGTGSLGSLGDRGAAPVTSSERLVDPNDLRKEDVAMELERVEDEEQWIKRSS 60
 DB 1 MGTGTGSLGSLGDRGAAPVTSSERLVDPNDLRKEDVAMELERVEDEEQWIKRSS 60
 QY 61 CNPLLOEPISAGGATAGTECKSVPCGWERVVKQNLFGKTAGRPVYVTSFGGLKFS 120
 DB 61 CNPLLOEPISAGGATAGTECKSVPCGWERVVKQNLFGKTAGRPVYVTSFGGLKFS 120

QY 121 KSSLANTYHKNGEISLKPEDFPTVLSKRGKSKRYKDCSMAALTSHLQNOGNSNMNIRT 180
 DB 121 KSSLANTYHKNGEISLKPEDFPTVLSKRGKSKRYKDCSMAALTSHLQNOGNSNMNIRT 180
 QY 181 RSKCKQVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKPKGKVTIILKG 240
 DB 181 RSKCKQVFMPPSSSELQESRGLSNFTSTHLLKEDGVDDVNRKVRKPKGKVTIILKG 240
 QY 241 IPIKTKKCKRSCSGFVQSDSKRESVCNKADABSEPAOKSOLDRTVCISDAGACGTL 300
 DB 241 IPIKTKKCKRSCSGFVQSDSKRESVCNKADABSEPAOKSOLDRTVCISDAGACGTL 300
 QY 301 SVTSEENLVKKKESLSSGNSPCSEOKTSGIINFCGAKOSEHNEKTEDTLESEBIC 360
 DB 301 SVTSEENLVKKKESLSSGNSPCSEOKTSGIINFCGAKOSEHNEKTEDTLESEBIC 360
 QY 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEKIFQEDTIPRTOIERRTSLYFSS 420
 DB 361 KVEVERKEHLHTDILKRGSEMDNNSPTRKDFTEKIFQEDTIPRTOIERRTSLYFSS 420
 QY 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLIIATITFLNRTSGMAIPVLMK 480
 DB 421 KYNKALSPRRKAFKKTTPRRSPNNLVOETLFHDPWKLIIATITFLNRTSGMAIPVLMK 480
 QY 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 DB 481 FLEKTPSAEAVARTADWRDVSLLKPLGLYDLRAKTIIVKPSDEYLTQWKYPIELHGIGKY 540
 QY 541 GNDSTRICVNMKQVHPEDHKLANKYHDMWMENHEKLSLS 580
 DB 541 GNDSTRICVNMKQVHPEDHKLANKYHDMWMENHEKLSLS 580

RESULT 4

AA44504
 ID AA44504 standard; protein; 565 AA.

AA44504;
 AC AA44504;

DT 27-MAR-2000 (first entry)
 XX

DE Human delta228-UV damage endonuclease.
 XX

XX Delta228-UVDE; ultraviolet damage endonuclease; GST signal peptide;
 KM glutathione-S-transferase signal peptide; uvei+ gene product;
 KM UV irradiation; DNA damage; UV radiation damage; photoproduct;
 KM abasic site; aplatinum diaduct; mismatched nucleotide pairing;
 KM nucleotide alkylation; skin cancer.
 XX

OS Homo sapiens.
 XX

XX WO963828-A1.
 XX

XX 16-DEC-1999.
 XX

XX 08-JUN-1999; 99WO-US012910.
 XX

XX 08-JUN-1998; 98US-0086521P.
 XX

XX 18-MAY-1999; 99US-0134752P.
 XX

XX (UYEM-) UNIV EMORY.
 XX

XX Doetsch PW, Kaur B, Avery AM;
 XX

XX MPI; 2000-116417/10.
 XX

XX A new truncated ultraviolet damage endonuclease for treatment of skin
 PT cancers.
 XX

XX Claim 16; Page 60; 133pp; English.
 XX

XX The present sequence is human delta228-UV damage endonuclease. Delta228-

PR	06-JUN-1997;	97US-0048949P.
PR	06-JUN-1997;	97US-0048962P.
PR	06-JUN-1997;	97US-0048963P.
PR	06-JUN-1997;	97US-0048964P.
PR	06-JUN-1997;	97US-0048965P.
PR	06-JUN-1997;	97US-0048970P.
PR	06-JUN-1997;	97US-0048971P.
PR	06-JUN-1997;	97US-0048972P.
PR	06-JUN-1997;	97US-0048974P.
PR	06-JUN-1997;	97US-0049019P.
PR	06-JUN-1997;	97US-0049020P.
PR	06-JUN-1997;	97US-0049373P.
PR	06-JUN-1997;	97US-0049374P.
PR	06-JUN-1997;	97US-0049375P.
PR	06-JUN-1997;	97US-0057584P.
PR	05-SEP-1997;	97US-0057627P.
PR	05-SEP-1997;	97US-0057628P.
PR	05-SEP-1997;	97US-0057629P.
PR	05-SEP-1997;	97US-0057632P.
PR	05-SEP-1997;	97US-0057634P.
PR	05-SEP-1997;	97US-0057635P.
PR	05-SEP-1997;	97US-0057642P.
PR	05-SEP-1997;	97US-0057643P.
PR	05-SEP-1997;	97US-0057644P.
PR	05-SEP-1997;	97US-0057645P.
PR	05-SEP-1997;	97US-0057646P.
PR	05-SEP-1997;	97US-0057647P.
PR	05-SEP-1997;	97US-0057648P.
PR	05-SEP-1997;	97US-0057649P.
PR	05-SEP-1997;	97US-0057650P.
PR	05-SEP-1997;	97US-0057651P.
PR	05-SEP-1997;	97US-0057654P.
PR	05-SEP-1997;	97US-0057661P.
PR	05-SEP-1997;	97US-0057662P.
PR	05-SEP-1997;	97US-0057666P.
PR	05-SEP-1997;	97US-0057667P.
PR	05-SEP-1997;	97US-0057668P.
PR	05-SEP-1997;	97US-0057760P.
PR	05-SEP-1997;	97US-0057761P.
PR	05-SEP-1997;	97US-0057762P.
PR	05-SEP-1997;	97US-0057763P.
PR	05-SEP-1997;	97US-0057764P.
PR	05-SEP-1997;	97US-0057765P.
PR	05-SEP-1997;	97US-0057769P.
PR	05-SEP-1997;	97US-0057770P.
PR	05-SEP-1997;	97US-0057771P.
PR	05-SEP-1997;	97US-0057774P.
PR	05-SEP-1997;	97US-0057775P.
PR	05-SEP-1997;	97US-0057776P.
PR	05-SEP-1997;	97US-0057777P.
PR	05-SEP-1997;	97US-0057778P.
PR	18-DEC-1997;	97US-0070923P.

cells comprising recombinant vectors containing the nucleic acid sequences are used for the recombinant production of the secreted proteins. The polynucleotide and amino acid sequences are useful for are useful for preventing, treating or ameliorating medical conditions e.g. by protein or gene therapy. Pathological conditions can be also diagnosed by determining the amount of the new polypeptides in a sample or by determining the presence of mutations in the new polynucleotides. Specific uses are described for each of the polynucleotides, based on which tissues they are most highly expressed in, and include developing products for the diagnosis or treatment of cancer, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, tumours, leukemias, diseases of the immune system, autoimmune diseases, hepatic and renal disease, lymphomas, inflammation, allergies, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, retinosis, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, diseases of testes, lung or thymus, digestive/endocrine disorders, infections and AIDS. The polypeptides are also useful for identifying their binding partners. The present sequence represents human secreted protein (see descriptor line for gene number and clone identification).

Query Match	40.3%	Score 1230;	DB 2;	Length 257;
Best Local Similarity	93.1%;	Pred. No. 3.3e-110;		
Matches 241; Conservative	3;	Mismatches 9;	Indels 6;	Gaps 1

QY 160 MAALTSHLONQSNNSNWNLTRSKCKKDVEMPPSSSELOESRGLSNFTSTHLLKEDEG 219

QY 220 VDDVNERKVRKPKGVTLKGIPIKTKKGCRCSCSGFVQSDSKRESVCNKDAESEPVA 279

[illegible]

Db 121 QKSQLDRTVCISDAGACGETLSTSEENSLVKKERSLSSGSNFCSEQKTSGIINKFCSA 180

Db 181 KDFHNEKYEDTFLESEBIGTKVEVERKEHHTDILKRGSEMDNNCSPTRKDF----- 235

QY 400 QEDTIPRTQIERRKTSLYF 418

ABB50468 standard; protein; 257 AA.

XX
XX
XX

DE Human secreted protein encoded by gene 168 SEQ ID NO:416.

KW dermatological; immunosuppressive; immunostimulant; autostatic; vascular; anti-angiogenic. ophthalmological:

KW antiparkinsonian; antitubercular; gene therapy; vaccine; immune disorder;
KW multiple sclerosis; systemic lupus erythematosus; HIV infection; cancer;
KW

KW Chaga's cardiomyopathy; coronary arteriosclerosis; angiogenic disorder;

KW Parkinson's disease; infectious disease; chromosome 3.

XX
OS
XX

BN WO200162891-A2.
XX
XX 30-AUG-2001.
XX
XX 21-FEB-2001; 2001WO-US005614.
XX
XX 24-FEB-2000; 2000US-0184836P.
XX 29-MAR-2000; 2000US-0193170P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Ni J, Ebner R, Lafleur DM, Moore PA, Olsen HS, Rosen CA;
PI Ruben SM, Soppet DR, Young PE, Shi Y, Florence KA, Mei Y;
PI Florence C, Hu J, Li Y, Kyaw H, Fischer CL, Ferris AM, Fan P,
PI Peng P, Andrews GA, Dillon PJ, Carter KC, Brewer LA, Yu G, Zeng Z,
PI Greene JM;
XX
XX WPI; 2001-625724/72.
DR N-PSDB; ABA83361.
XX
XX Nucleic acids encoding 207 human secreted polypeptides, useful for
PI preventing, diagnosing and/or treating, e.g. cancers, Parkinson's disease
PI and diabetic retinopathy.
XX
XX Claim 11; Page 1171-1172; 1533pp; English.
XX
XX ABB50301 to ABB51287 and ABA83194 to ABA83441 represent human secreted
CC proteins (I) and polynucleotide (II) sequences. (I) and (II) have various
CC activities based on the tissues and cells the genes are expressed in.
CC Example of these activities include: immunomodulatory; antisclerotic;
CC dermatological; immunosuppressive; anti-inflammatory; immunostimulant;
CC anti-HIV; cytostatic; cardiant; anti-angiogenic; ophthalmological;
CC neuroprotective; nootropic; anticonvulsant; antiallergens; vascular;
CC antiparkinsonian; antimicrobial; and vulnerary. (I) and (II) can be used
CC in gene therapy and vaccine production. (I) and (II) can be used in the
CC prevention, diagnosis and treatment of immune disorders (e.g. multiple
CC sclerosis, systemic lupus erythematosus and human immunodeficiency virus
CC (HIV) infections), hyperproliferative disorders (e.g. cancers and
CC Gaucher's disease), cardiovascular diseases (e.g. Schmitz syndrome,
CC Chaga's cardiomyopathy and coronary arteriosclerosis), angiogenic
CC disorders (e.g. corneal graft neovascularisation and diabetic
CC retinopathy), neurological disorders (e.g. Huntington's chorea,
CC Alzheimer's disease and Parkinson's disease), infectious diseases and/or
CC for promoting wound healing, regeneration and/or chemotaxis. ABA83185 to
CC ABA83193 and ABB50300 represent sequences used in the exemplification of
CC the present invention
XX
XX Sequence 257 AA;
SQ
Query Match 40.3%; Score 1230; DB 4; Length 257;
Best Local Similarity 93.1%; Pred. No. 3.3e-110;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
QY 160 MAALTSHLONQNNNSNMNRTSKCKKQVFMPPSSSELOEGRGSLNFTSTHLLKEDSG 219
DB 1 MAALTSHLONQNNNSNMNRTSKCKKQVFMPPSSSELOEGRGSLNFTSTHLLKEDSG 60
QY 220 VDVNPRKRRKPKGKTIKGIPIKTKKGRKSCGFPQSDSKREYCNKADASEEPYA 279
DB 61 VDVNPRKRRKPKGKTIKGIPIKTKKGRKSCGFPQSDSKREYCNKADASEEPYA 120
QY 280 QKSQDLRTVCISDAGACGETLSTVSEENSLVKKKERSLSSGSNFCSEQKTSGLINKFCGA 339
DB 121 QKSQDLRTVCISDAGACGETLSTVSEENSLVKKKERSLSSGSNFCSEQKTSGLINKFCGA 180
QY 340 KQSEHNKEKEDPTLESEEGTKVYVERKEHLTDILKXGSEMDNKCSTRKDFGEKTF 399
DB 181 KQSEHNKEKEDPTLESEEGTKVYVERKEHLTDILKXGSEMDNKCSTRKDFGEKTF 235
QY 400 QEDTTPRTQIERKKTLYF 418
DB 236 -EDTTPRNTDRKKENKPVF 253

RESULT 9
ABO44725
ID ABO44725 standard; protein; 257 AA.
XX
XX ABO44725;
XX
XX 02-OCT-2003 (first entry)
XX
XX Novel human secreted protein #168.
XX
XX Human; gene therapy; autoimmune disorder; multiple sclerosis; cancer;
XX systemic lupus erythematosus; haematopoietic cell disorder; allergy;
XX agammaglobulinaemia; ataxia telangiectasia; blood coagulation disorder;
XX althrombocytopenia; thrombocytopenia; graft-versus-host disease; arthritis;
XX inflammatory condition; ischaemia-reperfusion injury; infectious disease;
XX hyperproliferative disorder; purpura; viral infection; regeneration;
XX bacterial infection; ulcer; Alzheimer's disease.
XX
XX Homo sapiens.
XX
XX US2003065160-A1.
XX
XX 03-APR-2003.
XX
XX 07-DEC-2001; 2001US-00004860.
XX
XX 06-JUN-1997; 97US-0048875P.
XX 06-JUN-1997; 97US-0048876P.
XX 06-JUN-1997; 97US-0048877P.
XX 06-JUN-1997; 97US-0048878P.
XX 06-JUN-1997; 97US-0048879P.
XX 06-JUN-1997; 97US-0048880P.
XX 06-JUN-1997; 97US-0048881P.
XX 06-JUN-1997; 97US-0048882P.
XX 06-JUN-1997; 97US-0048883P.
XX 06-JUN-1997; 97US-0048884P.
XX 06-JUN-1997; 97US-0048885P.
XX 06-JUN-1997; 97US-0048892P.
XX 06-JUN-1997; 97US-0048893P.
XX 06-JUN-1997; 97US-0048894P.
XX 06-JUN-1997; 97US-0048895P.
XX 06-JUN-1997; 97US-0048896P.
XX 06-JUN-1997; 97US-0048897P.
XX 06-JUN-1997; 97US-0048898P.
XX 06-JUN-1997; 97US-0048899P.
XX 06-JUN-1997; 97US-0048900P.
XX 06-JUN-1997; 97US-0048901P.
XX 06-JUN-1997; 97US-0048915P.
XX 06-JUN-1997; 97US-0048916P.
XX 06-JUN-1997; 97US-0048917P.
XX 06-JUN-1997; 97US-0048949P.
XX 06-JUN-1997; 97US-0048962P.
XX 06-JUN-1997; 97US-0048963P.
XX 06-JUN-1997; 97US-0048964P.
XX 06-JUN-1997; 97US-0048970P.
XX 06-JUN-1997; 97US-0048971P.
XX 06-JUN-1997; 97US-0048972P.
XX 06-JUN-1997; 97US-0048974P.
XX 06-JUN-1997; 97US-0049019P.
XX 06-JUN-1997; 97US-0049020P.
XX 06-JUN-1997; 97US-0049373P.
XX 06-JUN-1997; 97US-0049374P.
XX 06-JUN-1997; 97US-0049375P.
XX 05-SEP-1997; 97US-0051584P.
XX 05-SEP-1997; 97US-0051627P.
XX 05-SEP-1997; 97US-0051628P.
XX 05-SEP-1997; 97US-0051629P.
XX 05-SEP-1997; 97US-0051634P.
XX 05-SEP-1997; 97US-0051635P.
XX 05-SEP-1997; 97US-0051642P.
XX 05-SEP-1997; 97US-0051643P.
XX 05-SEP-1997; 97US-0051644P.
XX 05-SEP-1997; 97US-0051645P.

PR 05-SEP-1997; 97US-0057646P.
PR 05-SEP-1997; 97US-0057647P.
PR 05-SEP-1997; 97US-0057648P.
PR 05-SEP-1997; 97US-0057649P.
PR 05-SEP-1997; 97US-0057650P.
PR 05-SEP-1997; 97US-0057651P.
PR 05-SEP-1997; 97US-0057654P.
PR 05-SEP-1997; 97US-0057661P.
PR 05-SEP-1997; 97US-0057662P.
PR 05-SEP-1997; 97US-0057663P.
PR 05-SEP-1997; 97US-0057664P.
PR 05-SEP-1997; 97US-0057665P.
PR 05-SEP-1997; 97US-0057669P.
PR 05-SEP-1997; 97US-0057700P.
PR 05-SEP-1997; 97US-0057717P.
PR 05-SEP-1997; 97US-0057718P.
PR 05-SEP-1997; 97US-0057775P.
PR 05-SEP-1997; 97US-0057776P.
PR 05-SEP-1997; 97US-0057777P.
PR 05-SEP-1997; 97US-0057778P.
PR 18-DEC-1997; 97US-0070923P.
PR 04-JUN-1998; 98MO-US011422.
PR 15-JUL-1998; 98US-0092921P.
PR 30-JUL-1998; 98US-0094657P.
PR 04-DEC-1998; 98US-00205258.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PI Young P, Greene JM, Ferrie AM, Ruben SM, Rosen CA, Hu J, J,
PI Olsen HS, Ebner R, Brewer LA, Moore PA, Shi Y, Florence C,
PI Florence K, Lafleur DW, Ni J, Fan P, Fischer CL, Soppet DR,
PI Li Y, Zeng Z, Kyaw H, Yu G, Feng P, Dillon PJ, Endress GA,
PI Carter KC,
XX
XX WPI; 2003-540804/51.
DR N-PSDB; ACH04862.
XX
XX
XX New isolated protein, useful for preparing a composition for diagnosing
XX or treating cancer, inflammatory, immune or infectious diseases.
PS Disclosure; SEQ ID NO 416; 172pp; English.
XX
XX The invention relates to an isolated HENAE80 protein. The protein is
XX useful for preparing a composition for diagnosing or treating autoimmune
XX disorders e.g. multiple sclerosis and systemic lupus erythematosus;
XX haematopoietic cell disorders e.g. agammaglobulinemia and ataxia
XX and telangiectasia; blood coagulation disorders e.g. afibrinogenemia and
XX thrombocytopenia; allergy; graft-versus-host disease; inflammatory
XX conditions e.g. ischaemia-reperfusion injury and arthritis;
XX hyperproliferative disorders e.g. cancer and purpura; infectious disease
XX e.g. viral infection and bacterial infection. The polynucleotide or
XX protein can be used to regenerate damaged tissue e.g. ulcers and
XX Alzheimer's disease. The present sequence represents the amino acid
XX sequence of a novel human secreted protein. Note: The sequence data for
XX this patent did not form part of the printed specification but was
XX obtained in electronic format directly from USPTO at
XX seqdata.uspto.gov/sequence.html?docID=20030065160
XX
XX Sequence 257 AA;
SQ
Query Match 40.3%; Score 1230; DB 6; Length 257;
Best Local Similarity 93.1%; Pred. No. 3.3e-110;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
OY 160 MAALTSHLQNSNNNNMLRTSRKCKKQVFMPPSSSSSELQSRGLSNTSTHLLKEDEG 219
1 MAALTSHLQNSNNNNMLRTSRKCKKQVFMPPSSSSSELQSRGLSNTSTHLLKEDEG 60

OY 220 VDDVNFRRKVRKPKGKVTLLKGIPIKTKKCGCRKSCSGFVQSDSKRESVCKNADESEPPVA 279
DB 61 VDDVNFRRKVRKPKGKVTLLKGIPIKTKKCGCRKSCSGFVQSDSKRESVCKNADESEPPVA 120
OY 280 QKSQLDRTVCISDAGACGFTLSVTSEENSLVKKKERSLSSGSNFCSEFOKTSGLINKCSA 339
DB 121 QKSQLDRTVCISDAGACGFTLSVTSEENSLVKKKERSLSSGSNFCSEFOKTSGLINKCSA 180
OY 340 KDSHNRKRYEDTFLSESEIGTKVVERKEHLHTDILKRGSEMDNCSPTRKDPTGEKIF 399
DB 181 KDSHNRKRYEDTFLSESEIGTKVVERKEHLHTDILKRGSEMDNCSPTRKDPTGEKIF 235
OY 400 QEDTTPRQIERKTSLYF 418
DB 236 -EDTTPRQIERKTSLYF 253
RESULT 10
ABO26205
ID ABO26205 standard; protein; 257 AA.
XX
XX ABO26205;
AC
XX
XX 10-SEP-2003 (first entry)
DT
XX
XX Human protein from novel secreted protein gene 168.
DE
XX
XX Human, secreted protein; precerebellin-like protein;
KM neurodegenerative disorder; behavioural disorder; Alzheimer's disease;
KM Parkinson's disease; Huntington's disease; schizophrenia; mania;
KM dementia; paranoia; psychosis; autism; immune disorder; infection;
KM inflammation; allergy; liver disorder; hepatoblastoma; jaundice;
KM hepatitis; immunological disorder; AIDS; leukaemia; rheumatoid arthritis;
KM sepsis; acne; psoriasis; cancer.
XX
XX Homo sapiens.
OS
XX
XX US6525174-B1.
PN
XX
XX 25-FEB-2003.
PD
XX
XX
XX 04-DEC-1998; 98US-00205258.
PP
XX
XX 06-JUN-1997; 97US-0048875P.
PR 06-JUN-1997; 97US-0048876P.
PR 06-JUN-1997; 97US-0048877P.
PR 06-JUN-1997; 97US-0048878P.
PR 06-JUN-1997; 97US-0048880P.
PR 06-JUN-1997; 97US-0048881P.
PR 06-JUN-1997; 97US-0048882P.
PR 06-JUN-1997; 97US-0048883P.
PR 06-JUN-1997; 97US-0048884P.
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PR 06-JUN-1997; 97US-0048887P.
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PR 06-JUN-1997; 97US-0048895P.
PR 06-JUN-1997; 97US-0048896P.
PR 06-JUN-1997; 97US-0048897P.
PR 06-JUN-1997; 97US-0048898P.
PR 06-JUN-1997; 97US-0048899P.
PR 06-JUN-1997; 97US-0048900P.
PR 06-JUN-1997; 97US-0048901P.
PR 06-JUN-1997; 97US-0048915P.
PR 06-JUN-1997; 97US-0048916P.
PR 06-JUN-1997; 97US-0048917P.
PR 06-JUN-1997; 97US-0048919P.
PR 06-JUN-1997; 97US-0048962P.
PR 06-JUN-1997; 97US-0048963P.
PR 06-JUN-1997; 97US-0048964P.
PR 06-JUN-1997; 97US-0048970P.
PR 06-JUN-1997; 97US-0048971P.

PR 06-JUN-1997; 97US-0048972P.
 PR 06-JUN-1997; 97US-0048974P.
 PR 06-JUN-1997; 97US-0049019P.
 PR 06-JUN-1997; 97US-0049020P.
 PR 06-JUN-1997; 97US-0049373P.
 PR 06-JUN-1997; 97US-0049374P.
 PR 06-JUN-1997; 97US-0049375P.
 PR 05-SEP-1997; 97US-0057584P.
 PR 05-SEP-1997; 97US-0057627P.
 PR 05-SEP-1997; 97US-0057628P.
 PR 05-SEP-1997; 97US-0057629P.
 PR 05-SEP-1997; 97US-0057634P.
 PR 05-SEP-1997; 97US-0057635P.
 PR 05-SEP-1997; 97US-0057642P.
 PR 05-SEP-1997; 97US-0057643P.
 PR 05-SEP-1997; 97US-0057644P.
 PR 05-SEP-1997; 97US-0057645P.
 PR 05-SEP-1997; 97US-0057646P.
 PR 05-SEP-1997; 97US-0057647P.
 PR 05-SEP-1997; 97US-0057648P.
 PR 05-SEP-1997; 97US-0057649P.
 PR 05-SEP-1997; 97US-0057650P.
 PR 05-SEP-1997; 97US-0057651P.
 PR 05-SEP-1997; 97US-0057654P.
 PR 05-SEP-1997; 97US-0057661P.
 PR 05-SEP-1997; 97US-0057662P.
 PR 05-SEP-1997; 97US-0057666P.
 PR 05-SEP-1997; 97US-0057667P.
 PR 05-SEP-1997; 97US-0057668P.
 PR 05-SEP-1997; 97US-0057760P.
 PR 05-SEP-1997; 97US-0057761P.
 PR 05-SEP-1997; 97US-0057762P.
 PR 05-SEP-1997; 97US-0057763P.
 PR 05-SEP-1997; 97US-0057765P.
 PR 05-SEP-1997; 97US-0057766P.
 PR 05-SEP-1997; 97US-0057769P.
 PR 05-SEP-1997; 97US-0057770P.
 PR 05-SEP-1997; 97US-0057771P.
 PR 05-SEP-1997; 97US-0057774P.
 PR 05-SEP-1997; 97US-0057775P.
 PR 05-SEP-1997; 97US-0057776P.
 PR 05-SEP-1997; 97US-0057777P.
 PR 05-SEP-1997; 97US-0057778P.
 PR 18-DEC-1997; 97US-0070923P.
 PR 04-JUN-1998; 98WO-US011422.
 PR 15-JUL-1998; 98US-0092921P.
 PR 30-JUL-1998; 98US-0094657P.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Young P, Greene JM, Ruben SM, Rosen CA, Hu J,
 PI Olsen HS, Ehner R, Brewer LA, Moore PA, Shi Y, Florence C;
 PI Florence K, Lafleur DM, Ni J, Fan P, Wei Y, Flascher CL, Soppet DR,
 PI Li Y, Zeng Z, Kyaw H, Yu G, Feng P, Dillon PJ, Endreass GA,
 PI Carter KC;
 XX
 DR MPI: 2003-511926/48.
 DR N-PsDB; ACD44672.
 XX
 PT New precerebellin-like protein, useful for diagnosing or treating
 PT neurodegenerative and behavioral disorders, immune disorders, liver
 PT disorders, and cancer.
 XX
 PS Disclosure; SEQ ID NO 416; 156bp; English.
 XX
 CC The invention relates to an isolated protein comprising amino acid
 CC residues 33-205 or 1-205 of a novel human secreted protein appearing as
 CC ABO26252. The protein is encoded by one of 238 disclosed cDNA sequences
 CC encoding 238 secreted proteins. ABO26252 is a precerebellin-like protein.
 CC Also included are a composition comprising the protein and a carrier and
 CC an isolated protein produced by expressing the protein cited above by a
 CC cell, and recovering the protein. The proteins are useful for diagnosing
 CC or treating neurodegenerative and behavioural disorders (e.g. Alzheimer's

CC disease, Parkinson's disease, Huntington's disease, schizophrenia, mania,
 CC dementia, paranoia, psychoses or autism), immune disorders (e.g.
 CC infection, inflammation, allergy), liver disorders (e.g. hepatoblastoma,
 CC jaundice, hepatitis), immunological disorders (e.g. AIDS, leukaemia,
 CC rheumatoid arthritis, sepsis, acne, psoriasis) and cancer. The present
 CC sequence is one of the 238 disclosed novel secreted proteins. Note: The
 CC sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from USPTO
 CC at: seqdata.uspto.gov/sequence.html?docid=6525174B1
 XX
 SQ Sequence 257 AA;
 Query Match 40.3%; Score 1230; DB 7; Length 257;
 Best Local Similarity 93.1%; Pred. No. 3,3e-110;
 Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
 QY 160 MAALTSHLONSNNSNNMLRTSRKCKDQFMPSSSSLSQSRGLSNFTSTHLIKEDBG 219
 DB 1 MAALTSHLONSNNSNNMLRTSRKCKDQFMPSSSSLSQSRGLSNFTSTHLIKEDBG 60
 QY 220 VDDVNFRRVRRPKKGVITLKGIPIKTKKGRKSCSGFVQSDSKRESVCNDAESEPVA 279
 DB 61 VDDVNFRRVRRPKKGVITLKGIPIKTKKGRKSCSGFVQSDSKRESVCNDAESEPVA 120
 QY 280 QKSQLDRTVCISDAGAGETLSVTSSENSLVKKKRSJSSGSNFCSEKQTSGLINKFCSA 339
 DB 121 QKSQLDRTVCISDAGAGETLSVTSSENSLVKKKRSJSSGSNFCSEKQTSGLINKFCSA 180
 QY 340 KDSHNEKEYEDTFLSESEIGTKVEVERKEHLHTDILKRGSEMDNCSPTRKDTGEKIF 399
 DB 161 KDSHNEKEYEDTFLSESEIGTKVEVERKEHLHTDILKRGSEMDNCSPTRKDTGEKIF 235
 QY 400 QEDTIPRQIERKRTSLYF 418
 DB 236 -EDTIPRMTDRKKNKPVF 253
 RESULT 11
 ADD89916
 ID ADD89916 standard; protein; 202 AA.
 XX
 AC ADD89916;
 XX
 XX 29-JAN-2004 (first entry)
 DT
 XX
 DE Human 5-methylcytosine DNA glycosylase N-terminal deletion mutant.
 XX
 KM Human; 5-methylcytosine DNA glycosylase; enzyme; Cpg; mutant; mutcin.
 XX
 OS Homo sapiens.
 XX
 PN WO2003078593-A2.
 XX
 PD 25-SEP-2003.
 XX
 PF 14-MAR-2003; 2003WO-US007933.
 XX
 PR 15-MAR-2002; 2002US-0364689P.
 XX
 PA (EPIC-) EPIGENOMICS AG.
 XX
 PT Lofton-Day CE, Day JK;
 PI
 XX
 DR MPI: 2003-779127/73.
 XX
 PT Labeling methylated or methylatable CpG sequences, useful e.g. for
 PT diagnostic detection of altered methylation, comprises replacing
 PT methylated cytosine by labeled cytosine.
 XX
 PS Claim 11; Page 71-72; 73pp; English.
 XX
 CC The present sequence is the protein sequence of an N-terminal deletion
 CC mutant of human 5-methylcytosine DNA glycosylase (5-MCG), in which amino

CC acid residue 1 corresponds to amino acid 379 of the full-length protein
CC ADD89906. 5-MCDG acts by cleaving glycosyl bonds at methylated CpG
CC sites of DNA, removing 5-methylcytosine from the DNA backbone as a free
CC base. The N-terminal deletion mutant shows enhanced deglycosylase
CC specificity towards CpG dinucleotide sequences. Human 5-MCDG can be used
CC in a claimed method for labelling CpG sequences corresponding to
CC methylated CpG sequences in an isolated DNA sample. The method comprises:
CC digesting the genomic DNA with a restriction endonuclease to produce
CC genomic DNA fragments; treating the genomic DNA fragments with 5-MCDG
CC such that one or more 5-methylcytosine bases are removed to produce
CC abasic genomic DNA fragments; and treating these abasic genomic DNA
CC fragments with base excision repair enzymes in the presence of labelled
CC dCTP such that 5-methylcytosine removed from the genomic DNA fragments by
CC 5-MCDG is replaced by labelled cytosine in the one or more corresponding
CC positions of the abasic genomic DNA fragments to produce labelled genomic
CC DNA fragments, so that specific labelling of CpG sequences corresponding
CC to methylated CpG sequences is achieved. The 5-MCDG is also used in a
CC claimed method for comparing CpG methylation status, extent or pattern
CC between or among reference and test genomic DNA samples, and in a claimed
CC method for labelling potentially-methylatable CpG sequences in CpG-
CC containing genomic DNA fragments. The methods are used to identify
CC methylated and/or potentially methylatable CpG dinucleotides in genomic
CC DNA, including comparison of methylation pattern between healthy and
CC diseased samples, for diagnosis.

SQ Sequence 202 AA;

Query Match 36.2%; Score 1106; DB 7; Length 202;

Best Local Similarity 100.0%; Pred. No. 2,4e-98; Mismatches 0; Gaps 0;

Matches 202; Conservative 0; Indels 0; Gaps 0;

QY 379 GSEMDNNGSPTRKDFTEGKIFQEDTTPRTQIERRTSLYFSSKYKKEALSPRRKAFKKW 438

DB 1 GSEMDNNGSPTRKDFTEGKIFQEDTTPRTQIERRTSLYFSSKYKKEALSPRRKAFKKW 60

QY 439 TPSPSPFNLVQETLPHDPMKLLIATIFLNRTSGKMAIPVLMKFLKYPSEAVARTADWRD 498

DB 61 TPSPSPFNLVQETLPHDPMKLLIATIFLNRTSGKMAIPVLMKFLKYPSEAVARTADWRD 120

QY 499 VSELKPRGLYDLRAKTIIVKFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEKQVHP 558

DB 121 VSELKPRGLYDLRAKTIIVKFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEKQVHP 180

QY 559 EDHKLNKYHDMWLNENHEKLSLS 580

DB 181 EDHKLNKYHDMWLNENHEKLSLS 202

RESULT 12

ADD89908 standard; protein; 416 AA.

AC ADD89908;

DT 29-JAN-2004 (first entry)

DE Chicken 5-methylcytosine DNA glycosylase.

KM Chicken; 5-methylcytosine DNA glycosylase; enzyme; CpG.

OS Gallus gallus.

PN W02003078593-A2.

PD 25-SEP-2003.

PF 14-MAR-2003; 2003WO-US007933.

PR 15-MAR-2002; 2002US-0364689P.

PA (EPIG-) EPIGENOMICS AG.

PI Lofton-Day CR, Day JK;

XX WPI; 2003-779127/73.
DR N-PSDB; ADD89907.
XX
XX Labeling methylated or methylatable CpG sequences, useful e.g. for
PT diagnostic detection of altered methylation, comprises replacing
PT methylated cytosine by labeled cytosine.

PS Claim 11; Page 56-58; 73pp; English.

XX The present sequence is the protein sequence of chicken 5-methylcytosine
CC DNA glycosylase (5-MCDG). The enzyme acts by cleaving glycosyl bonds at
CC methylated CpG sites of DNA, removing 5-methylcytosine from the DNA
CC backbone as a free base. Chicken 5-MCDG can be used in a claimed method
CC for labelling CpG sequences corresponding to methylated CpG sequences in
CC an isolated DNA sample. The method comprises: digesting the genomic DNA
CC with a restriction endonuclease to produce genomic DNA fragments;
CC treating the genomic DNA fragments with 5-MCDG such that one or more 5-
CC methylcytosine bases are removed to produce abasic genomic DNA fragments;
CC and treating these abasic genomic DNA fragments with base excision repair
CC enzymes in the presence of labelled dCTP such that 5-methylcytosine
CC removed from the genomic DNA fragments by 5-MCDG is replaced by labelled
CC cytosine in the one or more corresponding positions of the abasic genomic
CC DNA fragments to produce labelled genomic DNA fragments, so that specific
CC labelling of CpG sequences corresponding to methylated CpG sequences is
CC achieved. The 5-MCDG is also used in a claimed method for comparing CpG
CC methylation status, extent or pattern between or among reference and test
CC genomic DNA samples, and in a claimed method for labelling potentially-
CC methylatable CpG sequences in CpG-containing genomic DNA fragments. The
CC methods are used to identify methylated and/or potentially methylatable
CC CpG dinucleotides in genomic DNA, including comparison of methylation
CC pattern between healthy and diseased samples, for diagnosis.

SQ Sequence 416 AA;

Query Match 28.8%; Score 880.5; DB 7; Length 416;

Best Local Similarity 78.2%; Pred. No. 6.3e-76; Mismatches 21; Indels 4; Gaps 2;

Matches 161; Conservative 20; Mismatches 21; Indels 4; Gaps 2;

QY 378 RGSMDNNGS--PTRKDFTEGKIFQ--EDTTPRTQIERRTSLYFSSKYKKEALSPRRK 433

DB 210 RGSMDNNGS--PTRKDFTEGKIFQ--EDTTPRTQIERRTSLYFSSKYKKEALSPRRK 269

QY 434 AFKKMTPSPSPFNLVQETLPHDPMKLLIATIFLNRTSGKMAIPVLMKFLKYPSEAVART 493

DB 270 AFKKMTPSPSPFNLVQETLPHDPMKLLIATIFLNRTSGKMAIPVLMKFLKYPSEAVART 329

QY 494 ADMKRVSELKPRGLYDLRAKTIIVKFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEW 553

DB 330 ADMKRVSELKPRGLYDLRAKTIIVKFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVNEW 389

QY 554 KQVHPEDHKLNKYHDMWLNENHEKLSL 579

DB 390 KQVHPEDHKLNKYHDMWLNENHEKLSL 415

RESULT 13

ADD89917 standard; protein; 147 AA.

AC ADD89917;

DT 29-JAN-2004 (first entry)

DE Human 5-methylcytosine DNA glycosylase N-terminal deletion mutant.

KM Human; 5-methylcytosine DNA glycosylase; enzyme; CpG; mutant; mutein.

OS Homo sapiens.

PN W02003078593-A2.

PD 25-SEP-2003.

PI

Query Match 11.7%; Score 357; DB 5; Length 68;
Best Local Similarity 98.5%; Pred. No. 2.1e-26;
Matches 67; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 81 ECRKSVPCGWERVVKORLFGTLAGRFEDVYFISPGGLKFRSSSLANYLHKNGETSLKPED 140
Db 1 ECRKSVPCGWERVVKORLFGTLAGRFEDVYFISPGGLKFRSSSLANYLHKNGETSLKPED 60

Qy 141 FDFTVLSK 148
Db 61 FDFTVLSK 68

RESULT 15
ABBS1147
ID ABBS1147 standard; protein; 50 AA.
XX
AC ABBS1147;
XX
DT 07-FEB-2002 (first entry)
XX
DE Human secreted protein encoded by gene 168 SEQ ID NO:1100.
XX
KW Human; secreted protein; immunomodulatory; antisclerotic; anti-HIV;
KW dermatological; immunosuppressive; antiinflammatory; immunostimulant;
KW cytosolic; cardiac; vascular; anti-angiogenic; ophthalmological;
KW neuroprotective; noctropic; anticonvulsant; antialzheimers; vulnerary;
KW antiparkinsonian; antitremor; gene therapy; vaccine; immune disorder;
KW multiple sclerosis; systemic lupus erythematosus; HIV infection; cancer;
KW human immunodeficiency virus; hyperproliferative disorder; wound healing;
KW Gaucher's disease; cardiovascular disease; Schmitz syndrome; chemotaxis;
KW Chaga's cardiomyopathy; coronary arteriosclerosis; angioecic disorder;
KW corneal graft neovascularization; diabetic retinopathy; regeneration;
KW neurological disorder; Huntington's chorea; Alzheimer's disease;
KW Parkinson's disease; infectious disease; chromosome 3.
XX
XX Homo sapiens.
OS
XX WO200162891-A2.
PN
XX 30-AUG-2001.
PD
XX 21-FEB-2001; 2001WO-US005614.
XX
XX 24-FEB-2000; 2000US-0184836P.
PR 29-MAR-2000; 2000US-0193170P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PA
XX
XX Ni J, Ebner R, Lafleur DW, Moore PA, Olsen HS, Rosen CA;
PI Ruben SM, Soppet DR, Young PE, Shi Y, Florence KA, Wei Y;
PI Florence C, Hu J, Li Y, Kyaw H, Fischer CL, Ferrie AM, Fan P;
PI Feng P, Endress GA, Dillon PJ, Carter KC, Brewer LA, Yu G, Zeng Z;
PI Greene JM;
XX
XX WPI; 2001-625724/72.
DR
XX
XX Nucleic acids encoding 207 human secreted polypeptides, useful for
PT preventing, diagnosing and/or treating, e.g. cancers, Parkinson's disease
PT and diabetic retinopathy.
PT
XX
XX Disclosure; Page 366; 1533pp; English.
PS
XX
XX ABB50301 to ABB51287 and ABA83194 to ABA83441 represent human secreted
CC proteins (I) and polynucleotide (II) sequences. (I) and (II) have various
CC activities based on the tissues and cells the genes are expressed in.
CC Example of these activities include: immunomodulatory; antisclerotic;
CC dermatological; immunosuppressive; antiinflammatory; immunostimulant;
CC anti-HIV; cytosolic; cardiac; anti-angiogenic; ophthalmological;
CC neuroprotective; noctropic; anticonvulsant; antialzheimers; vascular;
CC antiparkinsonian; antitremor; and vulnerary. (I) and (II) can be used
CC in gene therapy and vaccine production. (I) and (II) can be used in the
CC prevention, diagnosis and treatment of immune disorders (e.g. multiple

CC sclerosis, systemic lupus erythematosus and human immunodeficiency virus
CC (HIV) infections), hyperproliferative disorders (e.g. cancers and
CC Gaucher's disease), cardiovascular diseases (e.g. Schmitz syndrome,
CC Chaga's cardiomyopathy and coronary arteriosclerosis), angiogenic
CC disorders (e.g. corneal graft neovascularisation and diabetic
CC retinopathy), neurological disorders (e.g. Huntington's chorea,
CC Alzheimer's disease and Parkinson's disease), infectious diseases and/or
CC for promoting wound healing, regeneration and/or chemotaxis. ABA83185 to
CC ABA83193 and ABB50300 represent sequences used in the exemplification of
CC the present invention
XX
XX Sequence 50 AA;
SQ

Query Match 8.3%; Score 254; DB 4; Length 50;
Best Local Similarity 98.0%; Pred. No. 1.3e-16;
Matches 49; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 110 FISPGGLKFRSSSLANYLHKNGETSLKPEDFDFTVLSKRGISRYDCS 159
Db 1 FISPGGLKFRSSSLANYLHKNGETSLKPEDFDFTVLSKRGISRYDCS 50

Search completed: August 22, 2005, 10:05:09
Job time : 172 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:09 ; Search time 163 Seconds
(without alignments)
1393.374 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLSLSLDGGAAPTV.....HKLNKTHDMLWENHEKLSLS 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1759131 seqs, 391586102 residues

Total number of hits satisfying chosen parameters: 1759131

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

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19: /cgn2_6/ptodata/1/pubppaa/US11_PUBCOMB.pep.*
20: /cgn2_6/ptodata/1/pubppaa/US11_NEW_PUB.pep.*
21: /cgn2_6/ptodata/1/pubppaa/US60_NEW_PUB.pep.*
22: /cgn2_6/ptodata/1/pubppaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3055	100.0	580	US-10-389-853-2	Sequence 2, Appli
2	3055	100.0	580	US-10-629-951-2	Sequence 2, Appli
3	3011	98.6	574	US-10-629-951-24	Sequence 24, Appl
4	1639	53.6	307	US-10-389-853-11	Sequence 11, Appl
5	1326	43.4	384	US-10-629-951-29	Sequence 29, Appl
6	1230	40.3	257	US-09-933-767-416	Sequence 416, App
7	1230	40.3	257	US-10-004-860-416	Sequence 416, App
8	1230	40.3	257	US-10-023-282-416	Sequence 416, App
9	1106	36.2	202	US-10-389-853-12	Sequence 12, Appl
10	880.5	28.8	416	US-10-389-853-4	Sequence 4, Appli
11	816	26.7	147	US-10-389-853-13	Sequence 13, Appli

12	697	22.8	126	15	US-10-629-951-32	Sequence 32, Appli
13	616	20.2	119	15	US-10-629-951-30	Sequence 30, Appli
14	443	14.5	85	15	US-10-629-951-37	Sequence 37, Appli
15	357	11.7	68	10	US-09-967-869A-7	Sequence 7, Appli
16	357	11.7	68	20	US-11-045-828-7	Sequence 7, Appli
17	330	10.8	443	16	US-10-739-930-5943	Sequence 5943, Ap
18	300.5	9.8	382	16	US-10-425-115-299452	Sequence 299452,
19	281	9.2	185	16	US-10-767-701-33975	Sequence 33975, A
20	254	8.3	50	10	US-09-933-767-1100	Sequence 1100, Ap
21	254	8.3	50	14	US-10-004-860-1100	Sequence 1100, Ap
22	254	8.3	50	14	US-10-023-282-1100	Sequence 286, App
23	209	6.8	476	16	US-10-408-765A-266	Sequence 7, Appli
24	208.5	6.8	560	16	US-10-475-681-7	Sequence 3, Appli
25	208.5	6.8	561	16	US-10-475-681-3	Sequence 31, Appli
26	192	6.3	132	15	US-10-629-951-31	Sequence 5, Appli
27	186	6.1	68	10	US-09-967-869A-5	Sequence 5, Appli
28	186	6.1	68	20	US-11-045-828-5	Sequence 5, Appli
29	152	5.0	105	16	US-10-437-963-148365	Sequence 148365,
30	145	4.7	1301	15	US-10-369-493-1644	Sequence 1644, Ap
31	145	4.7	3418	15	US-10-392-113-42	Sequence 42, Appli
32	145	4.7	3418	15	US-10-634-574-1	Sequence 1, Appli
33	145	4.7	3418	15	US-10-408-765A-178	Sequence 178, App
34	137	4.5	560	14	US-10-205-841-12	Sequence 12, Appli
35	137	4.5	1713	18	US-10-840-512-116	Sequence 116, App
36	136.5	4.5	409	16	US-10-425-115-250086	Sequence 250086,
37	136.5	4.5	1359	17	US-10-732-923-8708	Sequence 8708, Ap
38	135.5	4.4	282	10	US-09-967-869A-17	Sequence 17, Appli
39	135.5	4.4	282	20	US-11-045-828-17	Sequence 17, Appli
40	135.5	4.4	755	16	US-10-473-127-548	Sequence 548, App
41	135.5	4.4	755	16	US-10-473-127-550	Sequence 550, App
42	135.5	4.4	755	16	US-10-473-127-553	Sequence 553, App
43	133.5	4.4	1359	17	US-10-732-923-8707	Sequence 8707, Ap
44	132	4.3	578	9	US-09-821-835-2	Sequence 2, Appli
45	132	4.3	1819	15	US-10-335-977-7981	Sequence 7981, Ap

ALIGNMENTS

RESULT 1
US-10-389-853-2
Sequence 2, Appli
Publication No. US20030180779A1
GENERAL INFORMATION:
APPLICANT: Loflon-Day, Cathy E.
TITLE OF INVENTION: Diagnostic Methods Using 5-Methylcytosine DNA Glyco
FILE REFERENCE: 47675-36
CURRENT APPLICATION NUMBER: US/10/389, 853
CURRENT FILING DATE: 2003-03-14
PRIOR APPLICATION NUMBER: 60/364, 689
PRIOR FILING DATE: 2002-03-15
NUMBER OF SEQ ID NOS: 13
SOFTWARE: PatentIn version 3.1
SEQ ID NO 2
LENGTH: 580
TYPE: PRT
ORGANISM: Homo sapiens
US-10-389-853-2

Query Match	100.0%	Score 3055	DB 14	Length 580
Best Local Similarity	100.0%	Pred. No. 8.2e-236	Indels 0	Gaps 0
Matches 580	Conservative 0	Mismatches 0	Indels 0	Gaps 0
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DB	61	CNPLDPEPIASQFGATAGTCRKSVPCCGWERVYKORLFGKTAGRPDYVFIFPQGIKFRS	120	
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Db 181 RSKCKDVMPMPSSSSSELOESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVTLTKG 240
Qy 241 IPKTKKGCGRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGRTL 300
Db 241 IPKTKKGCGRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGRTL 300
Qy 301 SVTSEENSLVKKERSLSGSGNFCSGEOKTGIIINKFCSAKOSEHNEKXEDTFLSEBEGT 360
Db 301 SVTSEENSLVKKERSLSGSGNFCSGEOKTGIIINKFCSAKOSEHNEKXEDTFLSEBEGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNCSPTKDFTEGKIPOEDTIPRTQIERRTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNCSPTKDFTEGKIPOEDTIPRTQIERRTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIGKY 540
Db 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIGKY 540
Qy 541 GNDSYRIFCVNEMKQVAPBEDHKLNTKYHDMLENHEKLSLS 580
Db 541 GNDSYRIFCVNEMKQVAPBEDHKLNTKYHDMLENHEKLSLS 580
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RESULT 2

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US-10-629-951-2
; Sequence 2, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-2
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Query Match 100.0%; Score 3055; DB 15; Length 580;
Best Local Similarity 100.0%; Pred. No. 8,26-236;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 1 MGTTLGSLSLGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVEDEEQMMIKRSSE 60
Qy 61 CNPLQEPPIASAOFGATAGTECRKSVPCGWERVVKQRLFGKTAGRFDVYFISPOGLKRS 120
Db 61 CNPLQEPPIASAOFGATAGTECRKSVPCGWERVVKQRLFGKTAGRFDVYFISPOGLKRS 120
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Db 121 KSSLANTYLNKGETSLKPEDFDFTVLSKRGKISKRYKDCSMALTSHLQONOSNNMNLRT 180
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Db 121 KSSLANTYLNKGETSLKPEDFDFTVLSKRGKISKRYKDCSMALTSHLQONOSNNMNLRT 180
Qy 181 RSKCKDVMPMPSSSSSELOESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVTLTKG 240
Db 181 RSKCKDVMPMPSSSSSELOESRGLSNFTSTHLLKEDGVDDVNFRRKVRKPKGVTLTKG 240
Qy 241 IPKTKKGCGRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGRTL 300
Db 241 IPKTKKGCGRKSCSGFVQSDSKRESVCKNKADESEPAOKSOLDRTVCISDAGACGRTL 300
Qy 301 SVTSEENSLVKKERSLSGSGNFCSGEOKTGIIINKFCSAKOSEHNEKXEDTFLSEBEGT 360
Db 301 SVTSEENSLVKKERSLSGSGNFCSGEOKTGIIINKFCSAKOSEHNEKXEDTFLSEBEGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNCSPTKDFTEGKIPOEDTIPRTQIERRTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNCSPTKDFTEGKIPOEDTIPRTQIERRTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKMTPPRSPFNLVQETLPHDPKLLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIGKY 540
Db 481 FLEKYPBAEVAARTADWDVSELKPLGLYDLRAKTIYKFSDEYLTQWKYPILHGIGKY 540
Qy 541 GNDSYRIFCVNEMKQVAPBEDHKLNTKYHDMLENHEKLSLS 580
Db 541 GNDSYRIFCVNEMKQVAPBEDHKLNTKYHDMLENHEKLSLS 580
```

RESULT 3

```
US-10-629-951-24
; Sequence 24, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 574
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-24
```

```
Query Match 98.6%; Score 3011; DB 15; Length 574;
Best Local Similarity 99.0%; Pred. No. 2,76-232;
Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1;
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```
Qy 1 MGTTLGSLSLGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVEDEEQMMIKRSSE 60
Db 1 MGTTLGSLSLGDRGAAPVTYSSERLVDPDPNDLRKEDVAMELERVEDEEQMMIKRSSE 60
Qy 61 CNPLQEPPIASAOFGATAGTECRKSVPCGWERVVKQRLFGKTAGRFDVYFISPOGLKRS 120
Db 61 CNPLQEPPIASAOFGATAGTECRKSVPCGWERVVKQRLFGKTAGRFDVYFISPOGLKRS 120
Qy 121 KSSLANTYLNKGETSLKPEDFDFTVLSKRGKISKRYKDCSMALTSHLQONOSNNMNLRT 180
Db 121 KSSLANTYLNKGETSLKPEDFDFTVLSKRGKISKRYKDCSMALTSHLQONOSNNMNLRT 180
```

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QY 181 RSKCKDVMPSPSSSEILOSERGLSNFTSTHLLKEDSGVDVNRKTRKPKGKTTLLKG 240
DB 181 RSKCKDVMPSPSSSEILOSERGLSNFTSTHLLKEDSGVDVNRKTRKPKGKTTLLKG 240
QY 241 IPIKTKKGCRCSCGSPVSDSKRSVCKADAESEPVAKSQDLRTVCISDAGCGEYL 300
DB 241 IPIKTKKGCRCSCGSPVSDSKRSVCKADAESEPVAKSQDLRTVCISDAGCGEYL 300
QY 301 SVTSEENSIVKKKERSLSGSGNFCSEOKTSGIINKFCSAKDEHNEKXEDTFLSEBEIGT 360
DB 301 SVTSEENSIVKKKERSLSGSGNFCSEOKTSGIINKFCSAKDEHNEKXEDTFLSEBEIGT 360
QY 361 KVEVERKEHHTDILKRGSEMDNNSPTRKOPFTGEKIFQEDTTPRTQIERKTSLYFS 420
DB 361 KVEVERKEHHTDILKRGSEMDNNSPTRKOPFTGEKIFQEDTTPRTQIERKTSLYFS 414
QY 421 KYNKALSPRRKAPFKMTPPRSPFNLVOETLPHDPWKLITIFPLNRTSGMAIPVLMK 480
DB 421 KYNKALSPRRKAPFKMTPPRSPFNLVOETLPHDPWKLITIFPLNRTSGMAIPVLMK 474
QY 445 FLEKTPSAEVARATADWRDVSSEILKPLGLYDLRAKTIKFSDEYLTQWKYPIELHGIGKY 540
DB 445 FLEKTPSAEVARATADWRDVSSEILKPLGLYDLRAKTIKFSDEYLTQWKYPIELHGIGKY 534
QY 541 GNDYRIFCVNEMKQVHPEDHKLNTKYHDLWLNENHEKLSLS 580
DB 541 GNDYRIFCVNEMKQVHPEDHKLNTKYHDLWLNENHEKLSLS 574
QY 535 GNDYRIFCVNEMKQVHPEDHKLNTKYHDLWLNENHEKLSLS 574
DB 535 GNDYRIFCVNEMKQVHPEDHKLNTKYHDLWLNENHEKLSLS 574

RESULT 4
US-10-389-853-11
/ Sequence 11, Application US/10389853
/ Publication No. US2003018079A1
/ GENERAL INFORMATION:
/ APPLICANT: Lofton-Day, Cathy E.
/ TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco
/ FILE REFERENCE: 47675-36
/ CURRENT APPLICATION NUMBER: US/10/389,853
/ PRIOR FILING DATE: 2003-03-14
/ PRIOR APPLICATION NUMBER: 60/364,689
/ PRIOR FILING DATE: 2002-03-15
/ NUMBER OF SEQ ID NOS: 13
/ SOFTWARE: PatentIn version 3.1
/ SEQ ID NO 11
/ LENGTH: 307
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ FEATURE:
/ NAME/KEY: NON TER
/ LOCATION: (1)-(11)
/ OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
/ OTHER INFORMATION: no.1 corresponds to aa no.274 of hMBD4; mutant shows enhanced d
/ OTHER INFORMATION: eglycosylase specificity towards CpG dinucleotide sequences; see
/ OTHER INFORMATION: Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
US-10-389-853-11

Query Match 53.6%; Score 1639; DB 14; Length 307;
Best Local Similarity 100.0%; Pred. No.1e-122;
Matches 307; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 274 ESEPVAKSQDLRTVCISDAGCGEYLSVTSEENSIVKKKERSLSGSGNFCSEOKTSGI 333
DB 1 ESEPVAKSQDLRTVCISDAGCGEYLSVTSEENSIVKKKERSLSGSGNFCSEOKTSGI 60
QY 334 NKFCSAKDEHNEKXEDTFLSEBEIGTKVEVERKEHHTDILKRGSEMDNNSPTRKOP 393
DB 61 NKFCSAKDEHNEKXEDTFLSEBEIGTKVEVERKEHHTDILKRGSEMDNNSPTRKOP 120
QY 394 TGEKIFQEDTTPRTQIERKTSLYFSKYNKALSPRRKAPFKMTPPRSPFNLVOETLF 453
DB 121 TGEKIFQEDTTPRTQIERKTSLYFSKYNKALSPRRKAPFKMTPPRSPFNLVOETLF 180
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QY 454 HDPWKLITATIFLNRTSGMAIPVLMKFLKCYPSAEVARATADWRDVSSEILKPLGLYDLRA 513
DB 181 HDPWKLITATIFLNRTSGMAIPVLMKFLKCYPSAEVARATADWRDVSSEILKPLGLYDLRA 240
QY 514 KTIKFSDEYLTQWKYPIELHGIGKYGNDYRIFCVNEMKQVHPEDHKLNTKYHDLWLN 573
DB 241 KTIKFSDEYLTQWKYPIELHGIGKYGNDYRIFCVNEMKQVHPEDHKLNTKYHDLWLN 300
QY 574 HEKLSLS 580
DB 301 HEKLSLS 307

RESULT 5
US-10-629-951-29
/ Sequence 29, Application US/10629951
/ Publication No. US20040018550A1
/ GENERAL INFORMATION:
/ APPLICANT: Bellacosa, Alfonso
/ TITLE OF INVENTION: Methods for Detection of Transition
/ FILE REFERENCE: FCCC 96-21
/ CURRENT APPLICATION NUMBER: US/10/629,951
/ PRIOR FILING DATE: 2003-07-29
/ PRIOR APPLICATION NUMBER: US/09/629,222A
/ PRIOR FILING DATE: 2000-07-31
/ PRIOR APPLICATION NUMBER: 09/463,891
/ PRIOR FILING DATE: 2000-01-28
/ PRIOR APPLICATION NUMBER: PCT/US98/15828
/ PRIOR FILING DATE: 1998-07-28
/ PRIOR APPLICATION NUMBER: 60/053,936
/ PRIOR FILING DATE: 1997-07-28
/ SOFTWARE: FaatSEO for Windows Version 3.0
/ SEQ ID NO 29
/ LENGTH: 384
/ TYPE: PRT
/ ORGANISM: Mus musculus
US-10-629-951-29

Query Match 43.4%; Score 1326; DB 15; Length 384;
Best Local Similarity 54.3%; Pred. No.1.7e-97;
Matches 283; Conservative 29; Mismatches 71; Indels 138; Gaps 4;

QY 36 KEDVAMELERVGEDEDEQMMIKRSSSECNPLQEPISAOFGATAGTGCRCXVPCGERVVK 95
DB 1 KEDVAMELERVGEDEDEQMMIKRSSSECNPLQEPISAOFGATAGTGCRCXVPCGERVVK 57
QY 96 QRLFGTKGRPDVYVYISFQGLKFRSKSLANYLHNKGETSLKRPDPFTVLSKGIKARY 155
DB 58 QRLFGTKGRPDVYVYISFQGLKFRSKSLANYLHNKGETSLKRPDPFTVLSKGIKARY 117
QY 156 KDCSMAALTSHLQONSSNNMNLARTRSCKKQVFMPPSSSEILOSERGLSNFTSTHLLK 215
DB 118 KDCSMAALTSHLQONSSNNMNLARTRSCKKQVFMPPSSSEILOSERGLSNFTSTHLLK 177
QY 216 BDEGVDVNFRRKVRKPKGVITLLKGIPIKTKKGCRCSCGSPVSDSKRSVCKADAE 275
DB 178 BDEGVDVNFRRKVRKPKGVITLLKGIPIKTKKGCRCSCGSPVSDSKRSVCKADAE 228
QY 276 EPVAKSQDLRTVCISDAGCGEYLSVTSEENSIVKKKERSLSGSGNFCSEOKTSGI 335
DB 229 ----- 228
QY 336 FCSAKDEHNEKXEDTFLSEBEIGTKVEVERKEHHTDILKRGSEMDNNSPTRKOP 395
DB 229 ----- 228
QY 396 EKIQEDTTPRTQIERKTSLYFSKYNKALSPRRKAPFKMTPPRSPFNLVOETLF 455
DB 229 -----EKSIPQVKKRKTSLYFSKYNKALSPRRKAPFKMTPPRSPFNLVOETLF 283
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Db      1 MAALTSHLQNSNSNMNLRTRSKCKKDVMPSPSSSELOESRGLSNFTSTHLLKEDBG 60
Qy      220 VDDVNRKRVKRPKGVTKLIGIPIKTKKGCRCRSCGFPVQSDSKESVCKNKADESEPYA 279
Db      61 VDDVNRKRVKRPKGVTKLIGIPIKTKKGCRCRSCGFPVQSDSKESVCKNKADESEPYA 120
Qy      280 QKSQDLRTVCISDAGACGETTISVTSEHNSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 339
Db      121 QKSQDLRTVCISDAGACGETTISVTSEHNSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 180
Qy      340 KQSEHNKEKEDTFLSEEEIGTVEVVERKEHLHTDILKRGSEMDNNSCPTRKDFTGEKLF 399
Db      181 KQSEHNKEKEDTFLSEEEIGTVEVVERKEHLHTDILKRGSEMDNNSCPTRKDFT----- 235
Qy      400 QEDTIPRTQIERRKTSLYF 418
Db      236 -EDTIPRNTDRKKENKPVF 253

RESULT 7
US-10-004-860-416
; Sequence 416, Application US/10004860
; Publication No. US20030065160A1
; GENERAL INFORMATION:
; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/10/004,860
; CURRENT FILING DATE: 2001-12-07
; Prior Application removed - See File Wrapper or Palm
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 416
; LENGTH: 257
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SITE
; LOCATION: (100)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-10-004-860-416

Query Match      40.3%; Score 1230; DB 14; Length 257;
Best Local Similarity 93.1%; Pred. No. 4,6e-90;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

Qy      160 MAALTSHLQNSNSNMNLRTRSKCKKDVMPSPSSSELOESRGLSNFTSTHLLKEDBG 219
Db      1 MAALTSHLQNSNSNMNLRTRSKCKKDVMPSPSSSELOESRGLSNFTSTHLLKEDBG 60
Qy      220 VDDVNRKRVKRPKGVTKLIGIPIKTKKGCRCRSCGFPVQSDSKESVCKNKADESEPYA 279
Db      61 VDDVNRKRVKRPKGVTKLIGIPIKTKKGCRCRSCGFPVQSDSKESVCKNKADESEPYA 120
Qy      280 QKSQDLRTVCISDAGACGETTISVTSEHNSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 339
Db      121 QKSQDLRTVCISDAGACGETTISVTSEHNSLVKKKERSLSSGNSFCSEQKTSGLINKFCSA 180
Qy      340 KQSEHNKEKEDTFLSEEEIGTVEVVERKEHLHTDILKRGSEMDNNSCPTRKDFTGEKLF 399
Db      181 KQSEHNKEKEDTFLSEEEIGTVEVVERKEHLHTDILKRGSEMDNNSCPTRKDFT----- 235
Qy      400 QEDTIPRTQIERRKTSLYF 418
Db      236 -EDTIPRNTDRKKENKPVF 253

RESULT 8
US-10-023-282-416
; Sequence 416, Application US/10023282
; Publication No. US20030092893A1
; GENERAL INFORMATION:
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; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/10/023,282
; CURRENT FILING DATE: 2001-12-20
; EARLIER APPLICATION NUMBER: 09/205,258
; EARLIER FILING DATE: 1998-12-04
; EARLIER APPLICATION NUMBER: PCT/US98/11422
; EARLIER FILING DATE: 1998-06-04
; EARLIER APPLICATION NUMBER: 60/048,885
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,375
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,881
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,880
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,896
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,020
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,876
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,895
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,884
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,894
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,971
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,964
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,882
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,899
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,893
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,900
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,901
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,892
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,915
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,019
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,970
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,972
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,916
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,373
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,875
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,374
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,917
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,949
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,974
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,883
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,897
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,898
; EARLIER FILING DATE: 1997-06-06
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EARLIER APPLICATION NUMBER: 60/048,962
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,963
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,877
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,878
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/070,923
EARLIER FILING DATE: 1997-12-18
EARLIER APPLICATION NUMBER: 60/092,921
EARLIER FILING DATE: 1998-07-15
EARLIER APPLICATION NUMBER: 60/094,657
EARLIER FILING DATE: 1998-07-30
NUMBER OF SEQ ID NOS: 1227
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 416
LENGTH: 257
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: SITE
LOCATION: (100)
OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-10-023-282-416

Query Match 40.3%; Score 1230; DB 14; Length 257;
Best Local Similarity 93.1%; Pred. No. 4,6e-90;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

QY 160 MAALTSHTLQNSNNMNLRTSKCKKQVFMPPSSSSSELOBSRGISNFTSTHLLKEDG 219
DB 1 MAALTSHTLQNSNNMNLRTSKCKKQVFMPPSSSSSELOBSRGISNFTSTHLLKEDG 60
QY 220 VDDVFRKVRKPKGKVTLLKGP1PIKTKYKCKKSCSGFVQSDSKRESVYCNADAESEFVA 279
DB 61 VDDVFRKVRKPKGKVTLLKGP1PIKTKYKCKKSCSGFVQSDSKRESVYCNADAESEFVA 120
QY 280 QKSQDRVCTSDACGCEITLVSEENSLVKKKRSISGSGNFCSEKTSGLINKPCSA 339
DB 121 QKSQDRVCTSDACGCEITLVSEENSLVKKKRSISGSGNFCSEKTSGLINKPCSA 180
QY 340 KQSENEKEDTFLSESEIGTKVEVVERKEHLHTDILKRGSEMDNCSPTKDFTEKIF 399
DB 181 KQSENEKEDTFLSESEIGTKVEVVERKEHLHTDILKRGSEMDNCSPTKDFTEKIF 235
QY 400 QEDTTPRTQIERKTSLYF 418
DB 236 -EDTTPRTQIERKTSLYF 253

RESULT 9
US-10-389-853-12
Sequence 12, Application US/10389853
Publication No. US20030180779A1
GENERAL INFORMATION:
APPLICANT: Lofton-Day, Cathy E.
TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco
FILE REFERENCE: 47675-36
CURRENT APPLICATION NUMBER: US/10/389,853
CURRENT FILING DATE: 2003-03-14
PRIOR APPLICATION NUMBER: 60/364,689
PRIOR FILING DATE: 2002-03-15
NUMBER OF SEQ ID NOS: 13
SOFTWARE: PatentIn version 3.1
SEQ ID NO 12
LENGTH: 202
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: NON_TER
LOCATION: (1)..(1)

OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
OTHER INFORMATION: no. 1 corresponds to aa no. 379 of hMBD4 protein; mutant shows en
OTHER INFORMATION: hanced deglycosylase specificity towards CpG dinucleotide sequenc
OTHER INFORMATION: es; see Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
US-10-389-853-12

Query Match 36.2%; Score 1106; DB 14; Length 202;
Best Local Similarity 100.0%; Pred. No. 2,8e-80;
Matches 202; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 379 GSEMDNCSPTKDFTEKIFQEDTTPRTQIERKTSLYSSSKYNKALSPPRKAKKW 438
DB 1 GSEMDNCSPTKDFTEKIFQEDTTPRTQIERKTSLYSSSKYNKALSPPRKAKKW 60
QY 439 TTPRSPNLVOETLPHDPWKLITATIPLNRTSGMAIPVMKFLKXPASAEVARTADMRD 498
DB 61 TTPRSPNLVOETLPHDPWKLITATIPLNRTSGMAIPVMKFLKXPASAEVARTADMRD 120
QY 499 VSELKPLGLYDLRAKTIKVSDEYLTQWKYP1ELHIGIKYGNDSYRIFCVNEMKQVHP 558
DB 121 VSELKPLGLYDLRAKTIKVSDEYLTQWKYP1ELHIGIKYGNDSYRIFCVNEMKQVHP 180
QY 559 EDHKLKTYHDMLWENHEKLSLS 580
DB 181 EDHKLKTYHDMLWENHEKLSLS 202

RESULT 10
US-10-389-853-4
Sequence 4, Application US/10389853
Publication No. US20030180779A1
GENERAL INFORMATION:
APPLICANT: Lofton-Day, Cathy E.

TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco
FILE REFERENCE: 47675-36
CURRENT APPLICATION NUMBER: US/10/389,853
CURRENT FILING DATE: 2003-03-14
PRIOR APPLICATION NUMBER: 60/364,689
PRIOR FILING DATE: 2002-03-15
NUMBER OF SEQ ID NOS: 13
SOFTWARE: PatentIn version 3.1
SEQ ID NO 4
LENGTH: 416
TYPE: PRT
ORGANISM: Gallus gallus
US-10-389-853-4

Query Match 28.8%; Score 880.5; DB 14; Length 416;
Best Local Similarity 78.2%; Pred. No. 9,6e-62;
Matches 161; Conservative 20; Mismatches 21; Indels 4; Gaps 2;

QY 378 RGEEMDNCS--PTKDFTEKIFQ--EDTTPRTQIERKTSLYSSSKYNKALSPPRK 433
DB 210 RDSAADGVSWPSDKKSFYAVQAPRGTEBSAPRTQVDRRTSPYFSSKYKELSPPRK 269
QY 434 AFKKWTPRSPNLVOETLPHDPWKLITATIPLNRTSGMAIPVMKFLKXPASAEVART 493
DB 270 AFKKWTPRSPNLVOETLPHDPWKLITATIPLNRTSGMAIPVMKFLKXPASAEVART 329
QY 494 ADMWVSELKPLGLYDLRAKTIKVSDEYLTQWKYP1ELHIGIKYGNDSYRIFCVNEM 553
DB 330 ADMWVSELKPLGLYDLRAKTIKVSDEYLTQWKYP1ELHIGIKYGNDSYRIFCVNEM 389
QY 554 KQVHEDHKLKTYHDMLWENHEKLSLS 579
DB 390 KQVHEDHKLKTYHDMLWENHEKLSLS 415

RESULT 11
US-10-389-853-13
Sequence 13, Application US/10389853
Publication No. US20030180779A1

```

; GENERAL INFORMATION:
; APPLICANT: Lofton-Day, Cathy E.
; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco
; FILE REFERENCE: 47675-36
; CURRENT APPLICATION NUMBER: US/10/389,653
; PRIOR FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: 60/364,689
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 13
; LENGTH: 147
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: NON TER
; LOCATION: (1)..(1)
; OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
; OTHER INFORMATION: no. 1 corresponds to aa no. 434 of hMBD4 protein; mutant shows en
; OTHER INFORMATION: hanced deglycosylase specificity towards CpG dinucleotide sequenc
; OTHER INFORMATION: es; see Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
US-10-389-853-13

Query Match          26.7%; Score 816; DB 14; Length 147;
Best Local Similarity 100.0%; Pred. No. 3.1e-57;
Matches 147; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 444 AAKKTPPSPSPNLVQETLPHDPWKLITATITFLNRTSGKAIPLVMKFLKYPSPSAVART 493
    1 AAKKTPPSPSPNLVQETLPHDPWKLITATITFLNRTSGKAIPLVMKFLKYPSPSAVART 60
DB 494 ADMRDVSELLKPLGLYDLRAKTIYKFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVEM 553
    61 ADMRDVSELLKPLGLYDLRAKTIYKFSDEYLTQWKYPIELHIGIKYGNDSYRIFCVEM 120
QY 554 KQVHPEDHKLNKYHDMWLNHEKLSLS 580
    121 KQVHPEDHKLNKYHDMWLNHEKLSLS 147
DB 121 KQVHPEDHKLNKYHDMWLNHEKLSLS 147

RESULT 12
US-10-629-951-32
; Sequence 32, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: PCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaSTSeq for Windows Version 3.0
; SEQ ID NO: 32
; LENGTH: 126
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-32

Query Match          22.8%; Score 697; DB 15; Length 126;
Best Local Similarity 100.0%; Pred. No. 8.6e-48;
Matches 126; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 455 DPWKLIATITFLNRTSGKAIPLVMKFLKYPSPSAVARTADMWDVSELLKPLGLYDLRAK 514
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DB 1 DPWKLIATITFLNRTSGKAIPLVMKFLKYPSPSAVARTADMWDVSELLKPLGLYDLRAK 60
QY 515 TIVKRSDEYLTQWKYPIELHIGIKYGNDSYRIFCVEMKQVHPEDHKLNKYHDMWLNH 574
    61 TIVKRSDEYLTQWKYPIELHIGIKYGNDSYRIFCVEMKQVHPEDHKLNKYHDMWLNH 120
QY 575 EKLSLS 580
    121 EKLSLS 126
DB 121 EKLSLS 126

RESULT 13
US-10-629-951-30
; Sequence 30, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: PCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaSTSeq for Windows Version 3.0
; SEQ ID NO: 30
; LENGTH: 119
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-30

Query Match          20.2%; Score 616; DB 15; Length 119;
Best Local Similarity 100.0%; Pred. No. 2.5e-41;
Matches 119; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 36 KEDVAMELERVGEDEDEQMKIRSSCNPLDQEPISAGFGATGECRKSVPQGMERYVK 95
    1 KEDVAMELERVGEDEDEQMKIRSSCNPLDQEPISAGFGATGECRKSVPQGMERYVK 60
DB 1 KEDVAMELERVGEDEDEQMKIRSSCNPLDQEPISAGFGATGECRKSVPQGMERYVK 60
QY 96 ORLFKGTGRFPDVFYRISFGGLKFRKSSILANYLHNGETSLKPEDPFTVLSKRGIKSR 154
    61 ORLFKGTGRFPDVFYRISFGGLKFRKSSILANYLHNGETSLKPEDPFTVLSKRGIKSR 119
DB 61 ORLFKGTGRFPDVFYRISFGGLKFRKSSILANYLHNGETSLKPEDPFTVLSKRGIKSR 119

RESULT 14
US-10-629-951-37
; Sequence 37, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: PCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629,951
; PRIOR FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaSTSeq for Windows Version 3.0
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; SEQ ID NO 37
; LENGTH: 85
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-629-951-37

Query Match 14.5%; Score 443; DB 15; Length 85;
Best Local Similarity 100.0%; Pred. No. 1,1e-27;
Matches 85; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 69 IASAFGATAGTECKSVPCGWERVVKQRLFGKTAGRPDYVFISPOGLKFRSKSLANYL 128
Db 1 IASAFGATAGTECKSVPCGWERVVKQRLFGKTAGRPDYVFISPOGLKFRSKSLANYL 60

Qy 129 HKNGETSLKPEDPFTVLSKRGIKS 153
Db 61 HKNGETSLKPEDPFTVLSKRGIKS 85

RESULT 15

US-09-967-869A-7
; Sequence 7, Application US/09967869A
; Publication No. US2003008252A1
; GENERAL INFORMATION:
; APPLICANT: WOLFE, Alan P.
; APPLICANT: URNOV, Fyodor
; APPLICANT: LAI, Albert
; APPLICANT: RASCHKE, Eva
; TITLE OF INVENTION: MODULATION OF GENE EXPRESSION USING LOCALIZATION
; TITLE OF INVENTION: DOMAINS
; FILE REFERENCE: 8325-0019 / S19
; CURRENT APPLICATION NUMBER: US/09/967,869A
; CURRENT FILING DATE: 2001-09-28
; NUMBER OF SEQ ID NOS: 17
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 7
; LENGTH: 68
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: MBD4
US-09-967-869A-7

Query Match 11.7%; Score 357; DB 10; Length 68;
Best Local Similarity 98.5%; Pred. No. 6,2e-21;
Matches 67; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 81 ECRKSVPCGWERVVKQRLFGKTAGRPDYVFISPOGLKFRSKSLANYLHKNGETSLKPED 140
Db 1 ECRKSVPCGWERVVKQRLFGKTAGRPDYVFISPOGLKFRSKSLANYLHKNGETSLKPED 60

Qy 141 PDPFTVLSK 148
Db 61 PDPFTVLSK 68

Search completed: August 22, 2005, 10:12:39
Job time : 164 secs

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OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:09 ; Search time 43 Seconds
(without alignments)
1006.894 Million cell updates/sec

Title: US-10-629-951-2

Perfect score: 3055
Sequence: 1 MGTGLESLSLGDGGAAPT.....HKLNRYHDLWENHEKLSLS 580

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 513545 seqs, 74649064 residues

Total number of hits satisfying chosen parameters: 513545

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA:*
1: /cgn2_6/prodata/1/1aa/5A_COMB.pep:*
2: /cgn2_6/prodata/1/1aa/5B_COMB.pep:*
3: /cgn2_6/prodata/1/1aa/5A_COMB.pep:*
4: /cgn2_6/prodata/1/1aa/5B_COMB.pep:*
5: /cgn2_6/prodata/1/1aa/5A_COMB.pep:*
6: /cgn2_6/prodata/1/1aa/5B_COMB.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3055	100.0	580	3	US-09-327-984A-38
2	3055	100.0	580	4	US-09-629-222A-2
3	3055	100.0	580	4	US-09-657-013-56
4	3011	98.6	574	4	US-09-629-222A-24
5	1326	43.4	384	4	US-09-629-222A-29
6	1230	40.3	257	4	US-09-205-258-416
7	697	22.8	126	4	US-09-629-222A-32
8	616	20.2	119	4	US-09-629-222A-30
9	443	14.5	85	4	US-09-629-222A-37
10	254	8.3	50	4	US-09-205-258-1100
11	220.5	7.2	467	4	US-09-657-013-69
12	220.5	7.2	467	4	US-09-657-013-70
13	213.5	7.0	486	4	US-09-657-013-74
14	209	6.8	476	4	US-09-657-013-75
15	208.5	6.8	477	4	US-09-657-013-63
16	208.5	6.8	486	4	US-09-657-013-55
17	208.5	6.8	486	4	US-09-657-013-59
18	208.5	6.8	486	4	US-09-657-013-60
19	208.5	6.8	486	4	US-09-657-013-62
20	208.5	6.8	486	4	US-09-657-013-65
21	208.5	6.8	486	4	US-09-657-013-68
22	208.5	6.8	486	4	US-09-657-013-72
23	208.5	6.8	486	4	US-09-657-013-73
24	208.5	6.8	486	4	US-09-949-016-10209
25	202.5	6.6	345	4	US-09-657-013-112
26	201	6.6	482	4	US-09-657-013-66
27	198.5	6.5	484	4	US-09-657-013-58

28	198.5	6.5	484	4	US-09-657-013-61	Sequence 61, Appl
29	198.5	6.5	484	4	US-09-657-013-67	Sequence 67, Appl
30	198.5	6.5	484	4	US-09-657-013-71	Sequence 71, Appl
31	194	6.4	219	4	US-09-513-999C-6132	Sequence 6132, Ap
32	192	6.3	132	4	US-09-629-222A-31	Sequence 31, Appl
33	186	6.1	92	4	US-09-657-013-64	Sequence 64, Appl
34	146.5	4.8	1579	4	US-08-755-587-184	Sequence 184, App
35	145	4.7	3418	2	US-08-639-501-2	Sequence 2, Appl
36	145	4.7	3418	2	US-08-603-753D-4	Sequence 2, Appl
37	145	4.7	3418	3	US-09-044-946-2	Sequence 44, Appl
38	145	4.7	3418	3	US-08-755-587-44	Sequence 44, Appl
39	145	4.7	3418	3	US-09-044-908-2	Sequence 2, Appl
40	145	4.7	3418	3	US-09-099-753-4	Sequence 4, Appl
41	145	4.7	3418	3	US-08-986-106-4	Sequence 4, Appl
42	144	4.7	2329	3	US-08-755-587-16	Sequence 16, Appl
43	136.5	4.5	598	4	US-09-538-092-1083	Sequence 1083, Ap
44	131.5	4.3	291	4	US-09-657-013-48	Sequence 48, Appl
45	131.5	4.3	291	4	US-09-949-016-6438	Sequence 6438, Ap

ALIGNMENTS

RESULT 1									
US-09-327-984A-38									
Sequence 38, Application US/09327984A									
Patent No. 6368594									
GENERAL INFORMATION:									
APPLICANT: Doetsch, Paul W.									
APPLICANT: Kaur, Balveen									
APPLICANT: Avery, Angela M.									
TITLE OF INVENTION: Broad Specificity DNA Damage Endonuclease									
FILE REFERENCE: 25-98									
CURRENT APPLICATION NUMBER: US/09/327,984A									
PRIOR FILING DATE: 1999-06-08									
PRIOR APPLICATION NUMBER: US 60/088,521									
PRIOR FILING DATE: 1998-06-08									
PRIOR APPLICATION NUMBER: US 60/134,752									
PRIOR FILING DATE: 1999-05-18									
NUMBER OF SEQ ID NOS: 39									
SOFTWARE: PatentIn Ver. 2.0									
SEQ ID NO 38									
LENGTH: 580									
TYPE: PRT									
ORGANISM: Homo sapiens									
US-09-327-984A-38									
Query Match									
Best Local Similarity 100.0%; Score 3055; DB 3; Length 580;									
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;									
QY	1	MGTGLESLSLGDGGAAPT	VS	SERLVPPDNDLRKEDVAMELERVGEDEEQMIRSSB	60				
DB	1	MGTGLESLSLGDGGAAPT	VS	SERLVPPDNDLRKEDVAMELERVGEDEEQMIRSSB	60				
QY	61	CNPLOEPIASQFGATAGTECKRSVPCGWERVVQRLGKTAGRDVPYFISQGLKPS	120						
DB	61	CNPLOEPIASQFGATAGTECKRSVPCGWERVVQRLGKTAGRDVPYFISQGLKPS	120						
QY	121	KSLIANYLHKNGETSLSKPEDFPFVLSKRGISKRYDCSMALTSHLQNSNNNNLT	180						
DB	121	KSLIANYLHKNGETSLSKPEDFPFVLSKRGISKRYDCSMALTSHLQNSNNNNLT	180						
QY	181	RSKCKDVPMPSSSSELQESRGLSNFTSTHLLKEDGVDDVNFPRKVRPKGKVTILKG	240						
DB	181	RSKCKDVPMPSSSSELQESRGLSNFTSTHLLKEDGVDDVNFPRKVRPKGKVTILKG	240						
QY	241	IPKTKTKGCRKSGCFVSDSKREBVCNKAAESFPVAKQGLDPTVCISDAGAGETL	300						
DB	241	IPKTKTKGCRKSGCFVSDSKREBVCNKAAESFPVAKQGLDPTVCISDAGAGETL	300						
QY	301	SVTSENSLVKKKERLSGSGNFCSBQKTSGLINKFCSAKDSBHNKEYEDTFLSEIEIGT	360						
DB	301	SVTSENSLVKKKERLSGSGNFCSBQKTSGLINKFCSAKDSBHNKEYEDTFLSEIEIGT	360						

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Db 301 SVTSEENSLVKKERSLSGSGNFCSBQKTSGLINKFCASDSEHNEKYEDTFLESEBGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEEKIFQEDTIPRQIERRKTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEEKIFQEDTIPRQIERRKTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFHDPMKLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFHDPMKLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKPSAEVARTADWRDVSELKPLGLYDLRAKTIYKFSDEYLTQMKYPIELHGIGKY 540
Db 481 FLEKPSAEVARTADWRDVSELKPLGLYDLRAKTIYKFSDEYLTQMKYPIELHGIGKY 540
Qy 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKSLS 580
Db 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKSLS 580

RESULT 2
US-09-629-222A-2
; Sequence 2, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: PCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629, 222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053, 936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-629-222A-2

Query Match 100.0%; Score 3055; DB 4; Length 580;
Best Local Similarity 100.0%; Pred. No. 2.2e-292;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MGTGLESLSLGDGGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEBQMMIKRSSE 60
Db 1 MGTGLESLSLGDGGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEBQMMIKRSSE 60
Qy 61 CNPLLOEPIASAOGATAGTECRKSVPCGWERVVKQRLFGKTAGRPDYFISPGILKFRS 120
Db 61 CNPLLOEPIASAOGATAGTECRKSVPCGWERVVKQRLFGKTAGRPDYFISPGILKFRS 120
Qy 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLQONSNNMNLRT 180
Db 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLQONSNNMNLRT 180
Qy 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVITLKG 240
Db 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVITLKG 240
Qy 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQDLRTVCI SDAGACGETL 300
Db 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQDLRTVCI SDAGACGETL 300
Qy 301 SVTSEENSLVKKERSLSGSGNFCSBQKTSGLINKFCASDSEHNEKYEDTFLESEBGT 360
Db 301 SVTSEENSLVKKERSLSGSGNFCSBQKTSGLINKFCASDSEHNEKYEDTFLESEBGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEEKIFQEDTIPRQIERRKTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEEKIFQEDTIPRQIERRKTSLYFSS 420
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Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEEKIFQEDTIPRQIERRKTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFHDPMKLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFHDPMKLIATIFLNRTSGKMAIPVLMK 480
Qy 481 FLEKPSAEVARTADWRDVSELKPLGLYDLRAKTIYKFSDEYLTQMKYPIELHGIGKY 540
Db 481 FLEKPSAEVARTADWRDVSELKPLGLYDLRAKTIYKFSDEYLTQMKYPIELHGIGKY 540
Qy 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKSLS 580
Db 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKSLS 580

RESULT 3
US-09-657-013-56
; Sequence 56, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Treaf
; FILE REFERENCE: HO-P0189US1/03905371
; CURRENT APPLICATION NUMBER: US/09/657, 013
; PRIOR FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152, 778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: Patentn version 3.1
; SEQ ID NO 56
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Human
US-09-657-013-56

Query Match 100.0%; Score 3055; DB 4; Length 580;
Best Local Similarity 100.0%; Pred. No. 2.2e-292;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MGTGLESLSLGDGGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEBQMMIKRSSE 60
Db 1 MGTGLESLSLGDGGAAPVTYSSERLVDPDPNDLRKEDVAMELERVGEDEBQMMIKRSSE 60
Qy 61 CNPLLOEPIASAOGATAGTECRKSVPCGWERVVKQRLFGKTAGRPDYFISPGILKFRS 120
Db 61 CNPLLOEPIASAOGATAGTECRKSVPCGWERVVKQRLFGKTAGRPDYFISPGILKFRS 120
Qy 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLQONSNNMNLRT 180
Db 121 KKSILANTYHKNGETSLKPEDPFTVLSKRGIKSRKYKDCSMAALTSHLQONSNNMNLRT 180
Qy 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVITLKG 240
Db 181 RSKCKKQVFMPPSSSSSELOESRGLSNFTSTHLLKEDBGVDVNFVRKRPKGVITLKG 240
Qy 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQDLRTVCI SDAGACGETL 300
Db 241 IPIKTKKGRKSCSGFVQSDSKRESVCNKADASEBPVAKSQDLRTVCI SDAGACGETL 300
Qy 301 SVTSEENSLVKKERSLSGSGNFCSBQKTSGLINKFCASDSEHNEKYEDTFLESEBGT 360
Db 301 SVTSEENSLVKKERSLSGSGNFCSBQKTSGLINKFCASDSEHNEKYEDTFLESEBGT 360
Qy 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEEKIFQEDTIPRQIERRKTSLYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNNGSPTRKDFGEEKIFQEDTIPRQIERRKTSLYFSS 420
Qy 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFHDPMKLIATIFLNRTSGKMAIPVLMK 480
Db 421 KYNKEALSPRRKAFKKMTPPRSPFNLYOETLFHDPMKLIATIFLNRTSGKMAIPVLMK 480
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Db 421 KKNKALSPRRKAKKATPPRSPPNLVQETLFHPWKLIIATITLNTSGMAIPVLMK 480
Qy 481 FLEKPSAEVARTADMRDVSELKPLGLYDLRAKTIKPSDEYLTQKWKYPIELHIGIKY 540
Db 481 FLEKPSAEVARTADMRDVSELKPLGLYDLRAKTIKPSDEYLTQKWKYPIELHIGIKY 540
Qy 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKLSLS 580
Db 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKLSLS 580

RESULT 4
US-09-629-222A-24
; Sequence 24, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellco, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaastSeq for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 574
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-629-222A-24

Query Match          98.6%; Score 3011; DB 4; Length 574;
Best Local Similarity 99.0%; Pred. No. 4,7e-288;
Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1;

Qy 1 MGTGLESISLGDGCAAPVTTSSESLVPPDPNDLRKEDVAMEERGEDBEQMMIRSSSE 60
Db 1 MGTGLESISLGDGCAAPVTTSSESLVPPDPNDLRKEDVAMEERGEDBEQMMIRSSSE 60
Qy 61 CNPLQEPPLASAGFAGTAGTECKRSVPCGWERVVKRLPFGTAGRDVYFISFQGLKFRS 120
Db 61 CNPLQEPPLASAGFAGTAGTECKRSVPCGWERVVKRLPFGTAGRDVYFISFQGLKFRS 120
Qy 121 KSLIAYLHNKGETSLKPEDFDTVLSKGIKSRKYDCSMAALTSHLQNSNNNNMLRT 180
Db 121 KSLIAYLHNKGETSLKPEDFDTVLSKGIKSRKYDCSMAALTSHLQNSNNNNMLRT 180
Qy 121 KSLIAYLHNKGETSLKPEDFDTVLSKGIKSRKYDCSMAALTSHLQNSNNNNMLRT 180
Db 121 KSLIAYLHNKGETSLKPEDFDTVLSKGIKSRKYDCSMAALTSHLQNSNNNNMLRT 180
Qy 181 RSKCKDQVFMPPSSSELQESRGLSNFTSTHLLKDEGVDDVNFVRKPKKVTTLTKG 240
Db 181 RSKCKDQVFMPPSSSELQESRGLSNFTSTHLLKDEGVDDVNFVRKPKKVTTLTKG 240
Qy 241 IPKTKKGGKRSKCSGFSVQDSKREBYCNKADAESEPVAKOSQULDRVVCISDAGACETL 300
Db 241 IPKTKKGGKRSKCSGFSVQDSKREBYCNKADAESEPVAKOSQULDRVVCISDAGACETL 300
Qy 301 SVTSEENSILVKKERSLSSGSNFCSEQKTSIGIINKFCASADSEHNEKYEDTFLSEBEIGT 360
Db 301 SVTSEENSILVKKERSLSSGSNFCSEQKTSIGIINKFCASADSEHNEKYEDTFLSEBEIGT 360
Qy 361 KVEVERKEHLATDILKRGSEMDNCSPTKDTGEKIFQEDTTPRQIRAKTSLYFSS 420
Db 361 KVEVERKEHLATDILKRGSEMDNCSPTKDTGEKIFQEDTTPRQIRAKTSLYFSS 420
Qy 421 KKNKALSPRRKAKKATPPRSPPNLVQETLFHPWKLIIATITLNTSGMAIPVLMK 480
Db 421 KKNKALSPRRKAKKATPPRSPPNLVQETLFHPWKLIIATITLNTSGMAIPVLMK 480
Qy 481 FLEKPSAEVARTADMRDVSELKPLGLYDLRAKTIKPSDEYLTQKWKYPIELHIGIKY 540
Db 481 FLEKPSAEVARTADMRDVSELKPLGLYDLRAKTIKPSDEYLTQKWKYPIELHIGIKY 540
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Qy 481 FLEKPSAEVARTADMRDVSELKPLGLYDLRAKTIKPSDEYLTQKWKYPIELHIGIKY 540
Db 481 FLEKPSAEVARTADMRDVSELKPLGLYDLRAKTIKPSDEYLTQKWKYPIELHIGIKY 540
Qy 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKLSLS 580
Db 541 GNDSYRIFCVNEMKQVHPEDHKLNKYHDMWLNENHEKLSLS 580

RESULT 5
US-09-629-222A-29
; Sequence 29, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellco, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FaastSeq for Windows Version 3.0
; SEQ ID NO 29
; LENGTH: 384
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-629-222A-29

Query Match          43.4%; Score 1326; DB 4; Length 384;
Best Local Similarity 54.3%; Pred. No. 5,3e-122;
Matches 283; Conservative 29; Mismatches 71; Indels 138; Gaps 4;

Qy 36 KEDVAMELERVEDEBQMMIKSSSECNPLQEPPLASAGFAGTAGTECKRSVPCGWERVVK 95
Db 1 KEDVAMELERVEDEBQMMIKSSSECNPLQEPPLASAGFAGTAGTECKRSVPCGWERVVK 95
Qy 96 QRLFGKTAGRPVYFISFQGLKFRSKSLANTLHNKGETSLKPEDFDTVLSKGIKSR 155
Db 96 QRLFGKTAGRPVYFISFQGLKFRSKSLANTLHNKGETSLKPEDFDTVLSKGIKSR 155
Qy 58 QRLSGTAGKFDVYFISFQGLKFRSKSLANTLHNKGETSLKPEDFDTVLSKGIKSR 117
Db 58 QRLSGTAGKFDVYFISFQGLKFRSKSLANTLHNKGETSLKPEDFDTVLSKGIKSR 117
Qy 156 KDCSMAALTSHLQNSNNNNMLRTSKCKDQVFMPPSSSELQESRGLSNFTSTHLLTK 215
Db 156 KDCSMAALTSHLQNSNNNNMLRTSKCKDQVFMPPSSSELQESRGLSNFTSTHLLTK 215
Qy 118 KQISLAAALTSLOPNETDVSKQMLKTRSKKKTIVLPSTGTSSESSGSLSNSACTLLR 177
Db 118 KQISLAAALTSLOPNETDVSKQMLKTRSKKKTIVLPSTGTSSESSGSLSNSACTLLR 177
Qy 216 EDEGVDDVNFVRKPKKVTTLTKGIPKTKKGGKRSKCSGFSVQDSKREBYCNKADAE 275
Db 216 EDEGVDDVNFVRKPKKVTTLTKGIPKTKKGGKRSKCSGFSVQDSKREBYCNKADAE 275
Qy 178 EHRDIDVDSEKRSKSKRTVTLKGAQSKTKQCKKSLLESTQRRKAS 228
Db 178 EHRDIDVDSEKRSKSKRTVTLKGAQSKTKQCKKSLLESTQRRKAS 228
Qy 276 EPPAKOSQULDRVVCISDAGACETLSVTSEENSILVKKERSLSSGSNFCSEQKTSIGIINK 335
Db 276 EPPAKOSQULDRVVCISDAGACETLSVTSEENSILVKKERSLSSGSNFCSEQKTSIGIINK 335
Qy 229 ----- 228
Db 229 ----- 228
Qy 336 PCSAKDSEHNEKYEDTFLSEBEIGTVEVERKEHLATDILKRGSEMDNCSPTKDTFG 395
Db 336 PCSAKDSEHNEKYEDTFLSEBEIGTVEVERKEHLATDILKRGSEMDNCSPTKDTFG 395
Qy 229 ----- 228
Db 229 ----- 228
Qy 396 EKIQEDTTPRQIRAKTSLYFSSKYNKALSPRRKAKKATPPRSPPNLVQETLFHD 455
Db 396 EKIQEDTTPRQIRAKTSLYFSSKYNKALSPRRKAKKATPPRSPPNLVQETLFHD 455
Qy 229 ----- 228
Db 229 ----- 228
Qy 456 PWTGLIATITLNTSGMAIPVLMKLEKYPSAEVARTADMRDVSELKPLGLYDLRAKT 515
Db 456 PWTGLIATITLNTSGMAIPVLMKLEKYPSAEVARTADMRDVSELKPLGLYDLRAKT 515
Qy 284 PWTGLIATITLNTSGMAIPVLMKLEKYPSAEVARTADMRDVSELKPLGLYDLRAKT 343
Db 284 PWTGLIATITLNTSGMAIPVLMKLEKYPSAEVARTADMRDVSELKPLGLYDLRAKT 343
Qy 516 IYKPSDEYLTQKWKYPIELHIGIK-KYGNDSYRIFCVNEMKQ 555
Db 516 IYKPSDEYLTQKWKYPIELHIGIK-KYGNDSYRIFCVNEMKQ 555
Qy 344 IYKPSDEYLTQKWKYPIELHIGIK-KYGNDSYRIFCVNEMKQ 384
Db 344 IYKPSDEYLTQKWKYPIELHIGIK-KYGNDSYRIFCVNEMKQ 384
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RESULT 6
US-09-205-258-416
; Sequence 416, Application US/09205258
; Patent No. 6525174
; GENERAL INFORMATION:
; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/09/205,258
; EARLIER FILING DATE: 1998-12-04
; EARLIER APPLICATION NUMBER: PCT/US98/11422
; EARLIER FILING DATE: 1998-06-04
; EARLIER APPLICATION NUMBER: 60/048,885
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,375
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,881
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,880
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,896
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,020
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,876
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,895
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,884
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,894
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,971
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,964
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,882
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,899
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,893
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,900
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,901
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,892
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,915
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,019
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,970
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,972
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,916
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,373
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,875
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,374
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,917
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,949
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,974
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,883

EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,897
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,898
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,962
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,963
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,877
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,878
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/070,923
; EARLIER FILING DATE: 1997-12-18
; EARLIER APPLICATION NUMBER: 60/092,921
; EARLIER FILING DATE: 1998-07-15
; EARLIER APPLICATION NUMBER: 60/094,657
; EARLIER FILING DATE: 1998-07-30
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 416
; LENGTH: 257
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SITE
; LOCATION: (100)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-09-205-258-416

Query Match 40.3%; Score 1230; DB 4; Length 257;
Best Local Similarity 93.1%; Pred. No. 8.2e-113;
Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

QY 160 MAALTSHLQNSNNNNLRTSKCKKQVMPSSSELQBSRGLSNFTSTHLLKDEG 219
DB 1 MAALTSHLQNSNNNNLRTSKCKKQVMPSSSELQBSRGLSNFTSTHLLKDEG 60

QY 220 VDDVNFRRKVRKPKKKTITLIGIPKTKKGCRCSCSGFVSDSKRBSVCNKADSESPVA 279
DB 61 VDDVNFRRKVRKPKKKTITLIGIPKTKKGCRCSCSGFVSDSKRBSVCNKADSESPVA 120

QY 280 QKSOLDRTVCISDPAAGETLSTVSEENSLVKKKERLSGSGNFCSBEQKTSGLINKFCSA 339
DB 121 QKSOLDRTVCISDPAAGETLSTVSEENSLVKKKERLSGSGNFCSBEQKTSGLINKFCSA 180

QY 340 KDSHNEKYEDTFLSESEIGTVKVEVERKSHLHTDILKRGSEMDNNCSPTRKDTGKIF 399
DB 181 KDSHNEKYEDTFLSESEIGTVKVEVERKSHLHTDILKRGSEMDNNCSPTRKDTGKIF 235

QY 400 QEDTIPRTQIERKTSLYF 418
DB 236 -EDTIPRTQIERKTSLYF 253

RESULT 7
US-09-629-222A-32
; Sequence 32, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; EARLIER FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28

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CURRENT FILING DATE: 2000-07-31
PRIOR APPLICATION NUMBER: 09/463,891
PRIOR FILING DATE: 2000-01-28
PRIOR APPLICATION NUMBER: PCT/US98/15828
PRIOR FILING DATE: 1998-07-28
PRIOR APPLICATION NUMBER: 60/053,936
PRIOR FILING DATE: 1997-07-28
NUMBER OF SEQ ID NOS: 73
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 37
LENGTH: 85
TYPE: PRT
ORGANISM: Homo sapiens
US-09-629-222A-37

Query Match      14.5%; Score 443; DB 4; Length 85;
Best Local Similarity 100.0%; Pred. No. 6.3e-36;
Matches      85; Conservative    0; Mismatches    0; Indels    0; Gaps    0

Cy      69 IASAFQFAGTACTEERKSPVPCGMEWYVQRLEGGTAGRPDYVFISPGKLPRSKSLANTL 128
Db      1 IASAQFGTAATETCRKSVPCCGMERVYKQRLFGKTAGRDPYVFISQGLRFSKSLANTL 60

Cy      129 HKNGETSLKPEDPFTVLRSKGIGS 153
Db      61 HKNGETSLKPEDPFTVLRSKGIGS 85

RESULT 10
US-09-205-258-1100
Sequence 1100, Application US/09205258
Patent No. 6525174
GENERAL INFORMATION:
APPLICANT: Young et al.
TITLE OF INVENTION: 207 Human Secreted Proteins
FILE REFERENCE: P2007P1
CURRENT APPLICATION NUMBER: US/09/205,258
CURRENT FILING DATE: 1998-12-04
EARLIER APPLICATION NUMBER: PCT/US98/11422
EARLIER FILING DATE: 1998-06-04
EARLIER APPLICATION NUMBER: 60/048,885
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/049,375
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,881
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,880
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,896
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/049,020
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,876
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,895
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,884
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,894
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,971
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,964
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,882
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,899
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,893
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,900
EARLIER FILING DATE: 1997-06-06
EARLIER APPLICATION NUMBER: 60/048,901

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; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,892
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,915
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,019
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,970
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,972
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,916
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,373
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,875
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,374
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,917
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,949
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,974
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,883
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,897
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,898
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,962
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,963
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,877
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,878
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/070,923
; EARLIER FILING DATE: 1997-12-18
; EARLIER APPLICATION NUMBER: 60/092,921
; EARLIER FILING DATE: 1998-07-15
; EARLIER APPLICATION NUMBER: 60/094,657
; EARLIER FILING DATE: 1998-07-30
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1100
; LENGTH: 50
; TYPE: prt
; ORGANISM: Homo sapiens
US-09-205-258-1100

Query Match      8.3%; Score 254; DB 4; Length 50;
Best Local Similarity 98.0%; Pred. No. 1,2e-17;
Matches 49; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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Qy      110 FTSPPGLKFRSKSLANYLHKGSTSLKPEDFTVLSSKRGIKSKYKDCS 159
Db      1 FSSPGLKFRSKSLANYLHKGSTSLKPEDFTVLSSKRGIKSKYKDCS 50
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```
RESULT 11
US-09-657-013-69
; Sequence 69, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of identifying Mutations in a Methyl-CPG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
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```
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; CURRENT FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 69
; LENGTH: 467
; TYPE: prt
; ORGANISM: Pro9
US-09-657-013-69

Query Match      7.2%; Score 220.5; DB 4; Length 467;
Best Local Similarity 25.7%; Pred. No. 1e-12;
Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
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Qy      22 SSERLVDPDPNDLKEDVAMELER--VGDEDEOMATKRS--SECNPFLQEPFLASAOFGA 76
Db      49 SSEH-QGEPEADEGKADMSSESAEENLAVPESSASPKORRVRVIDRGPMYEDP----- 99
Qy      77 TAGTECKRSVPCGMEYVVKQRLFGKTAGRPDYVFISPGILKPRSKSLANYLHKNGETSL 136
Db      100 -----TLPEGWTRLKKQKSGRSAGKFDVYLINENGAFRSKVELIAYFOKVGDTSL 151
Qy      137 KPEDFTVLSSKRGIKSKRYDCSMALTSILOQSNNSNNLRTSRCKKDV--FMPPSS 194
Db      152 DPNDFTVL--TGKSPERRQG-----KPKKTKAPKSSVSGRGKPKSIIKKVPRV 204
Qy      195 SSELQESRGLSNFTSTHLIK----EDGEVDVNERKV-----RKPKQVTILKGI 242
Db      205 SEGQVAKRYIEK--SPGLLVMPYSGTKEASDATTSSQVLVILIRGGRKRS--TDPSAAP 262
Qy      243 IKTTKGCRKSCSGFVSDSKRESVCKADABEAPVAKQSDLRVTGISAAGCETLSV 302
Db      263 KKRKRKPSNVSLAAAEAAKKKAI--KESSTKPLIE-----TVLPKRRKTRRTISV 313
Qy      303 TSEE-----NSLVKK-----KERSLSSGNSFCSEKTSGLINKFCSAKDESHNEKYE 349
Db      314 DVKDTLPPEPLTVPIEKVMGQGNPAKSPESRSTGSKITGTGPKELQDQNNNNNNNNH 373
Qy      350 DTFLESEELGTQVVERKEHL 371
Db      374 HHSESKASATSPPEPTSKONI 395
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```
RESULT 12
US-09-657-013-70
; Sequence 70, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; CURRENT FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 70
; LENGTH: 467
; TYPE: prt
; ORGANISM: Pro9
US-09-657-013-70

Query Match      7.2%; Score 220.5; DB 4; Length 467;
Best Local Similarity 25.7%; Pred. No. 1e-12;
Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
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QY 22 SSERLVPDPNDLRKEDVAMELER--VGEDEEQMIKRS--SECNPLLOEPIASAOFGA 76
DB 49 SSEH-QPGEPADEGKADMSSESAEENILAVBESSASPKORSVLRDRGPMVEDP----- 99
QY 77 TAGTCRKSVPQGMERVVKQRLFGKTAGRPDVYFISPOGLKFRSKSLANYLHNKGETSL 136
DB 100 -----TLPEGMWTRKCLKQKRSKRSAGFPDYLLINPGKARSTVELLAFQKVGDTSL 151
QY 137 KPEPDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSMNLRTSKCKQDV--FMPSS 194
DB 152 DPNDPDFTV-TGRGSPSRREQ-----KQPKPKAPKPSVSGRGKGRPKGSIKXKYPVK 204
QY 195 SSELOESRGLSNFTSTHLLK-----EDEGVDVNFVKY-----RKPKGVTLIKGIP 242
DB 205 SEGVQVKRIEYK-SPGKLLVKWPGYSGTKEASDATTSQVLVIKRGGRKXSE-TDPSAP 262
QY 243 IKTKKCGKRSKSGVQSDSKRESVCNKADASEPVAOKSOLDRTVCISDAGAGETLSV 302
DB 263 KRGGRKPSNVSLAAAAAEAAKKKAI--KESIKPLLE-----TVLPKRRKRTETLSV 313
QY 303 TSEE-----NSLVKK-----KERSLSSGNSFCSEQKTSGLINKFCSAKDSEHNEKYE 349
DB 314 DYKDTIKPEPLTPVLEKWKQGNPAKSPBSRSTEGSPKIKTGLPKKELQOHNNHHHHHH 373
QY 350 DTFLESEIGTKVEVERKEHL 371
DB 374 HHSESKASATSPPEPETSMDNI 395

RESULT 13
US-09-657-013-74
; Sequence 74, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of identifying mutations in a Methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; PRIOR FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: Patentin version 3.1
; LENGTH: 486
; TYPE: PRT
; ORGANISM: Human
US-09-657-013-74

Query Match 7.0%; Score 213.5; DB 4; Length 486;
Best Local Similarity 24.4%; Pred. No. 5.3e-12;
Matches 106; Conservative 53; Mismatches 147; Indels 129; Gaps 18;

QY 37 EDVAMELERVGEDEEQMIKRSSECNPLLOEPIASAOFG-----ATACT 80
DB 22 KQKPLFKKVKKDKKEKEGKHEPVQPSAHHSABEAEKAEKTSBGSGSAPAVPEKASAP 81
QY 81 ECKRSV-----PCGWERVVKQRLFGKTAGRPDVYFISPOGLKFRSKSLANY 127
DB 82 KQKRSIIRDRGPMYDDPTLPEGMWTRKCLKQKRSKRSAGKTDVYLINQGAFRSKVELIAY 141
QY 128 LHNKGETSLKPEPDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSMNLRTSKCKQDV 187
DB 142 FEKVGDTSIDPNDPDFTV-TGRGSPSR-----REQKPPK- 175
QY 188 VFMPPSSSELQSRGL-----SNFTSTHLLKDEGVDVNFVKVRKPKGVTLIKGIPK 244
DB 176 ---PKSPKAPGTRGRGRPKSGGTTTPKKAATSEGVQVK--RVLEKSPGK--LLVMPP- 226

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QY 245 KTKGCKRSKSGFVQSDS-----KRESVCNKADASEPVAOKSOLDRTVCISDAGAGETL 300
DB 227 QTSFGKAEAGGATTSQVMYIKRGRKRAEADPOAIPKRR-----GKPKSV 276
QY 301 SVTSEENSLVKKERSLSSGNSFCSEQKTSGLINKFCSAKDSEHNEKEDTFLESEIGT 360
DB 277 AAAAEE--AKKK--AVKSSIRSVOETVLPIK-----RKTRET-----V 312
QY 361 KVEVERKEHLHTDIL--KRGSEMDNNSPTRKDPFGKTLFOEDTTPRTQIERKTSLYF 418
DB 313 SIEVKEVVKPLVSTLAEKSGKGLTKCKSPGR-----SKESPGR----- 354
QY 419 SSKTKKELSPRRK 433
DB 355 ---SSASPPKKE 365

RESULT 14
US-09-657-013-75
; Sequence 75, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of identifying mutations in a Methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; PRIOR FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: Patentin version 3.1
; LENGTH: 476
; TYPE: PRT
; ORGANISM: Human
US-09-657-013-75

Query Match 6.8%; Score 209; DB 4; Length 476;
Best Local Similarity 24.0%; Pred. No. 1.4e-11;
Matches 104; Conservative 53; Mismatches 149; Indels 128; Gaps 17;

QY 37 EDVAMELERVGEDEEQMIKRSSECNPLLOEPIASAOFG-----ATACT 81
DB 13 KQKPLFKKVKKDKKEKEGKHEPVQPSAHHSABEAEKAEKTSBGSGSARLCEASAP 72
QY 82 CRKSV-----PCGWERVVKQRLFGKTAGRPDVYFISPOGLKFRSKSLANYL 128
DB 73 QRSIIRDRGPMYDDPTLPEGMWTRKCLKQKRSKRSAGKTDVYLINQGAFRSKVELIAY 132
QY 129 HKNKGETSLKPEPDFTVLSKRGIKSRKYDCSMAALTSHLQNSNNSMNLRTSKCKQDV 188
DB 133 EKVGDTSIDPNDPDFTV-TGRGSPSR-----REQKPPK-- 165
QY 169 FMPSSSELQSRGL-----SNFTSTHLLKDEGVDVNFVKVRKPKGVTLIKGIPK 245
DB 166 ---PKSPKAPGTRGRGRPKSGGTTTPKKAATSEGVQVK--RVLEKSPGK--LLVMPP- 217
QY 246 TKGGRKSCSGFVQSDS-----KRESVCNKADASEPVAOKSOLDRTVCISDAGAGETLS 301
DB 218 TSPGKAEAGGATTSQVMYIKRGRKRAEADPOAIPKRR-----GRKGSVYA 267
QY 302 VTSSENSLVKKERSLSSGNSFCSEQKTSGLINKFCSAKDSEHNEKEDTFLESEIGT 361
DB 268 AAAAEE--AKKK--AVKSSIRSVOETVLPIK-----RKTRET-----V 303
QY 362 VEVVERKEHLHTDIL--KRGSEMDNNSPTRKDPFGKTLFOEDTTPRTQIERKTSLYF 419
DB 304 IVEVKEVVKPLVSTLAEKSGKGLTKCKSPGR-----SKESPGR----- 344

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QY      420 SKYNKEALSPRRK 433
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Db      345 ---SSSASSPPKKE 355
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RESULT 15

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US-09-657-013-63
; Sequence 63, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CpG-Binding Domain
; TITLE OF INVENTION: Containing Gene or Protein in Neurodevelopmental Disease and Tre
; FILE REFERENCE: HO-P01893US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657,013
; CURRENT FILING DATE: 2000-09-07
; PRIOR APPLICATION NUMBER: US 60/152,778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 63
; LENGTH: 477
; TYPE: prt
; ORGANISM: Human
US-09-657-013-63

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Query Match	6.8%	Score 208.5;	DB 4;	Length 477;
Best Local Similarity	23.9%	Pred. No. 1.6e-11;		
Matches 104; Conservative	53;	Mismatches 149;	Indels 129;	Gaps 17;

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QY      37 EDVAMELEAVGDEBQMIRKRSCECNPLLOEPIASNOFG-----ATAGT 80
Db      13 KKKPLKFKKKVKKDKKEKEGKEHPVOPSAHHSAPBAGCALETSEGSGBAPAVEKASP 72
QY      81 ECRKSV-----PCGWERVVKORLFEKTAGREFDVYETISFOGLFRSKSLANY 127
Db      73 KQRRSIIIRRGMYDDPPLPEGGWTRLKQKRSGRSAGKTDVYLINQGAFRSKVELIAY 132
QY      128 LHKNGETSLKPEDFDPTVLSKRGIKSRYYDCSMALLTSHLOQNSNNMNLFRRSCKKD 187
Db      133 FEKVADPTSLDPPNDFPETY-TGRSPSR-----REQKPPKK- 166
QY      188 VEMPPSSSESLQESRCL--SNFTSTHLLKEDBGVDVNFVRKRRPKGVTLLIKCIPIK 244
Db      167 ----FKSPRAPGTGKRGKRPKSGTTRPRKAATSEGVQYK--KVLKSPKK--LLVCMFP- 217
QY      245 KTKYKCGRKSCSGPFVOSDS---KRESVCKNKADESEPPVAOKSOLDRTVCISDAGAGETLU 300
Db      218 QTSPPGAKABGCGATSTQVMVIMKRPKRKKAEADPOAIPKKR-----GRKGGSVY 267
QY      301 SYTSEENSLVKKKERSLSSGNSNFCSPKQTSGLINRKCCKAKDSHNEKYEPTPLESBEIGT 360
Db      268 AAAAAABAKKAAVESSIR-----SVOETVLPIKK-----RKTRET-----V 303
QY      361 KVEAVERKEHHTDIL--KRGSMDNNCSPTPKDFTGEXKIFOEDTIPRTOIEHRKSTSLYF 418
Db      304 SIEVEAVVPLVSTLGEKSGKGLKTKCSBPGRK-----SKESSPPKGR----- 345
QY      419 SSKYNKEALSPPRK 433
Db      346 ----SSSASSPPKKE 356

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Search completed: August 22, 2005, 10:09:50
Job time : 44 secs